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# Chromosome 8-14 translocation in a non-African Burkitt's lymphoma with leukemic conversion. 

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#### Abstract

A specific chromosome translocation, $\mathrm{t}(8 \mathrm{q}-; 14 \mathrm{q}+)$, was observed in a 43-year-old female with non-African Burkitt's lymphoma in which leukemic conversion had occurred. The chromosome studies used cells from ascites. The ascites was apparently the result of a primary tumor involving the ovaries and contained $68 \%$ of lymphoma cells. The frequent occurrence of abnormalities related to chromosomes 1, 8 and 14 in African and non-African Burkitt's lymphomas was emphasized.


KEYWORDS: chromosome translocation, non-African Burkitt's lymphoma, mic conversion

# CHROMOSOME 8-14 TRANSLOCATION IN A NON-AFRICAN 

 BURKITT'S LYMPHOMA WITH LEUKEMIC CONVERSIONKanji Miyamoto, Jiro Sato, Koichi Kitajima*, Shunkichi Hiraki* Kohsuke Mori* and Toshio Tanaka**<br>Division of Pathology, Cancer Institute; * Department of Medicine (2nd Clinic); and<br>** Pathology Section, Central Laboratories, Okayama University Medical School, Okayama 700, Japan<br>Recived August 77, 1981


#### Abstract

A specific chromosome translocation, $\mathrm{t}(8 \mathrm{q}-; 14 \mathrm{q}+)$, was observed in a 43 -year-old female with non-African Burkitt's lymphoma in which leukemic conversion had occurred. The chromosome studies used cells from ascites. The ascites was apparently the result of a primary tumor involving the ovaries and contained $68 \%$ of lymphoma cells. The frequent occurrence of abnormalities related to chromosomes 1, 8 and 14 in African and non-African Burkitt's lymphomas was emphasized.


Key words : chromosome translocation, non-African Burkitt's lymphoma, leukemic conversion.

A specific translocation, $t(8 q-\cdots ; 14 q+)$, has been observed in African and non-African Burkitt lymphomas (1-3). The significance of this marker chromosome is of comparable importance to the original observation of the Philadelphia chromosome ( $\mathrm{Ph}^{1}$ ) rearrangement in chronic myelogenous leukemia (4). Recently, we reported on the Burkitt lymphoma of two Japanese, in whom a $14 q+$ marker chromosome was found $(5,6)$. This paper briefly reports another case of $t(8 q-$; $14 q+$ ) translocation in a Japanese Burkitt lymphoma with negative Epstein-Barr virus (EBV).

The patient was a 43-year-old female of single status. Towards the end of March, 1981, she developed abnormal genital bleeding and abdominal fullness; enlarged bilateral ovaries and ascites were detected at a gynecological examination. Peripheral blood (WBC: $10,400 / \mu \mathrm{l}$ ) on admission in the middle of April, contained $23 \%$ of immature cells, and the bone marrow biopsy demonstrated the typical "starry sky" effect (Fig. 1), indicating the leukemic coversion of Burkitt lymphoma. Ascites was apparently due to a primary tumor involving the ovaries, and contained approximately $68 \%$ of lymphoma cells (Fig. 2). These cells proved to be negative for EBV-determined nuclear antigen. Marker analysis showed that $90 \%$ of cells were positive for surface IgM.



Chromosomes were studied on cells from this ascites. The cells were incubated for 24 h in RPMI 1640 medium with $10 \%$ fetal calf serum and at $37^{\circ} \mathrm{C}$ in a humidified $5 \% \mathrm{CO}_{2}$ atmosphere. Mitotic cells were accumulated with Colcemid $(0.5 \mu \mathrm{~g} / \mathrm{ml})$, treated in a hypotonic solution of 75 mM KCl for 13 min , and fixed in a methanol-acetic acid $(3 ; 1)$ mixture. Chromosome preparations were stained with conventional Giemsa solution and analyzed using the Qbanding technique. Lymphoma cells had a modol number of 46 chromosomes; all the banded metaphases showed an identical karyotype, i.e., 46, XX, dir dup (lq) (pter - q32::q12 - q31::q32 - qter), $\mathrm{t}(8 ; 14)(\mathrm{q} 24 ; \mathrm{q} 32)$ (Fig. 3).

In the present study, we demonstrated a $t(8 q-; 14 q+)$ translocation in a Japanese adult with non-African Burkitt lymphoma. In addition, there was partial duplication of the long arm of chromosome No. 1. Douglass et al. (7) described a similar duplication of the long arms of chromosome No. 1. as well as the $14 q+$ marker in non-African Burkitt lymphoma. As shown by Slater et al. (8), $1 \mathrm{q}+$ rearrangement appears to play an important role in the evolution of the malignant cell population in lymphoproliferative disorders. The present case along with the two cases reported previously by us $(5,6)$ strongly suggests that abnormalities related to chromosomes 1,8 and 14 can be a frequent occurrence in Burkitt lymphoma in Japanese patients, in the same way as is seen in that of African, North American and European subjects.

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Fig. 1. Bone marrow biopsy showing the typical "starry sky" effect. H.E., $\times 200$.
Fig. 2. Lymphoma cells from ascites showing deeply stained scant cytoplasm with multiple uniform intracytoplasmic vacuoles. May-Grünwald-Giemsa, $\times 400$.

Fig. 3. Karyotype from ascites: $46, \mathrm{XX}$, dir dup ( $\mathbf{l q}$ ) (pter $-\mathrm{q} 32:: \mathrm{q} 12-\mathrm{q} 31:: \mathrm{q} 32-\mathrm{qter}$ ), $\mathrm{t}(8 ; 14)\left(\mathrm{q} 24 ; \mathrm{q}^{32}\right)$.
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