Impairment of T Cell Development in δEF1 Mutant Mice

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Summary

Using the method of gene targeting in mouse embryonic stem cells, regulatory function of δEF1, a zinc finger and homeodomain-containing transcription factor, was investigated in vivo by generating the δEF1 mutant mice. The mutated allele of δEF1 produced a truncated form of the δEF1 protein lacking a zinc finger cluster proximal to COOH terminus. The homozygous δEF1 mutant mice had poorly developed thymi with no distinction of cortex and medulla. Analysis of the mutant thymocyte showed reduction of the total cell number by two orders of magnitude accompanying the impaired thymocyte development. The early stage intrathymic c-kit+ T precursor cells were largely depleted. The following thymocyte development also seemed to be affected as assessed by the distorted composition of CD4- or CD8-expressing cells. The mutant thymocyte showed elevated α4 integrin expression, which might be related to the T cell defect in the mutant mice. In the peripheral lymph node tissue of the mutant mice, the CD4+CD8+ single positive cells were significantly reduced relative to CD4+CD8- single positive cells. In contrast to T cells, other hematopoietic lineages appeared to be normal. The data indicated that δEF1 is involved in regulation of T cell development at multiple stages.

Recent progress in our understanding of the T cell development clarified a major developmental pathway in thymus at cellular level: T cell precursors that originate from hematopoietic stem cells located in fetal liver and in adult bone marrow migrate and colonize in thymus. Starting from the CD4-CD8- double negative (DN) stage, thymocytes begin to rearrange their TCR genes and express CD3, a TCR coreceptor molecule, then proceed to the CD4+CD8- double positive (DP) stage. The DP thymocytes go through positive and negative selections depending on the specificity of the TCR. Finally, the CD4+CD8- or CD4-CD8+ single positive (SP) mature thymocytes are produced, and these immunocompetent cells migrate out and populate the peripheral lymphoid organs (1).

Some of these steps have been assigned to specific genes, and mutant mice of such genes produced by gene targeting have contributed greatly in defining each regulatory step of T cell development (2). However, it is obvious that more knowledge of genetic regulation is required to understand cellular events in T cell development. The δEF1 mutant mice to be reported in this paper has a novel phenotype:

1 Abbreviations used in this paper: DN, double negative; DP, double positive; d.p.c., days post coitum; FITC, fluorescein isothiocyanate; HBS, HEPES-buffered saline; SP, single positive.

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one of the major phenotype of both homozygous mutant mice was impaired of thymus development: severe hypopcellularity in thymus without clear distinction of cortex and medulla. Since Null–LacZ homozygous mutant mice are perinatally lethal with skeletal defects (to be published elsewhere), while ~20% of the ΔC-fin homozygous mutant mice were born alive and grown up to adulthood, we analyzed the lymphoid tissues in detail using the surviving young adult. ΔC-fin homozygous mutant mice. Here we describe the generation and analysis of ΔC-fin mutant mice and demonstrate that the defect of the thymus was ascribed to depletion of T precursor cells and to aberration of intrathymic development of T cells.

Materials and Methods

Mice. C57BL/6 and ICR mice were purchased from Japan SLC Inc. (Shizuoka, Japan) or CLEA Japan Inc. (Tokyo, Japan). All analyses were conducted under specific pathogen-free conditions. Constructions of Targeting Vector. Cloning and structural analysis of mouse ΔEF1 has been described (9). The targeting vector (see Fig. 2 A) was constructed as follows. A 0.8-kb SalI–Sall fragment containing a Sau3AI–SalI genomic fragment of the exon 6 sequence (see Fig. 2 A) and a 12-bp SalI–Sall linker adapter sequence, which is derived from an EMBL3 cloning vector, was subcloned into the Sall site of pBlueScript II to give pSS. The Sall and Xbal sites at the 5’ end of the inserted genomic fragment of pSS were inactivated by digesting with Xbal partially with Sall, blunt-ending by fill-in, and self-ligation. A Xbal linker carrying stop codons in all three frames (CTAGTCTAGACTAG) was inserted in the remaining Sall site at the 3’ end of the insert to have pSSstop. A XhoI–KpnI fragment of pSTNeoB (10) containing neo sequences was inserted in the XhoI–KpnI site of pSSstop, to have pSSstopNEO. In parallel, the 5.4-kb SalI–Apal genomic fragment, immediately 3' of the Sau3AI–Sall fragment was once cloned into the pBlueScript II, and regenerated by digesting with Asp718. The resulting fragment was blunt-ended, digested with Sall and cloned into the Sall–EcoRI V site of DST-A vector (11), generating pSADTA plasmid. The pSSStopNEO was digested with the Sall and Xbal sites at the 5’ end of the insert to have pSSstop. A XhoI–KpnI fragment of pSTNeoB (10) containing neo sequence was inserted in the XhoI–KpnI site of the pSSstop, to have pSSstopNEO. In parallel, the 5.4-kb SalI–Apal genomic fragment, immediately 3’ of the Sau3AI–Sall fragment was once cloned into the pBlueScript II, and regenerated by digesting with Asp718. The resulting fragment was blunt-ended, digested with Sall and cloned into the Sall–EcoRI V site of DST-A vector (11), generating pSADTA plasmid. The pSSStopNEO was digested with the Asp718, blunt-ended and digested with NotI. The resulting Asp718 (blunt-ended by Klenow)-NotI fragment was cloned into the Sall (blunt-ended by Klenow)-NotI sites of the pSADTA, generating a final targeting vector. The vector plasmid was linearized with NotI and used for electroporation. The expected targeted gene product lacks the COOH-terminal zinc finger clustes which have been shown to be essential for the DNA binding of ΔEF1 protein (12). The neo element has a promoter but lacks the termination and poly(A) addition signals, so that the neo is expressed only when poly(A) addition signal is supplied by recombination with a host gene. A DT-A cassette (11) was placed in the 3' end of the linearized vector for the negative selection against integration into nonhomologous genes.

Gene Targeting. E14 embryonic stem cells were electroporated and selected in the presence of G418 as described previously (13). Homologous recombinants were screened using Southern blot analysis, and obtained at a frequency of one in 107 electroporated cells. The ES cells carrying the mutated ΔEF1 allele were injected into blastocysts from C57BL/6 X C3H) F1 female mated with C57BL/6 male, and transferred to ICR pseudopregnant recipient mice. Resulting male chimeras were bred to ICR female mice to have heterozygous mice. The heterozygous mice were crossed with ICR or separately with C57BL/6 to keep the heterozygous pedigrees and intercrossed to generate homozygous mutant mice. Back-crossing to C57BL/6 has been done for six generations to date to obtain the mutant animals in C57BL/6 genetic background. Genotypes were determined by PCR analysis of DNA from ear punching (14) or by Southern blot analysis of tail DNA.

Southern and Northern Blot Analysis. Total DNA from the ES cells, the yolk sacs and the tails were isolated as described previously (15). The probe used in Southern blot analysis for identification of homologous recombination and genotyping was a 1.5-kb EcoRI–XbaI genomic fragment at 0.5 kb upstream of exon 6 (shown in Fig. 2 A). Total RNA of 12.5 d.p.c. embryos were prepared by a single step isolation procedure (16). 5 μg of the total RNA was separated by electrophoresis in a 1% formaldehyde agarose gel and blotted to a Hybond N (Amersham, Buckinghamshire, England) nylon membrane. The filter was hybridized with the 32P-labeled 2.5-kb EcoRI fragment of mouse ΔEF1 cDNA containing the sequence from exon 3 to exon 8 (9). Expression of the ΔEF1 ΔNAs in cultured Cells. A cDNA coding for ΔC-fin ΔEF1 protein was constructed by inserting the XbaI linker with stop codons in the Sall site of exon 6 of full-length cDNA in the same way as targeting vector construction. The full-length and ΔC-fin ΔEF1 ΔNAs were inserted into the NotI site of pCDM8 (17) and transfected to COS-7 cells by lipofection as described (18). An antiserum against N-proximal Portion of ΔEF1. An antiserum against human homologue of ΔEF1, AR E6, was used to detect the N-proximal region of mouse ΔEF1. HpaI–HaeIII 1041-2498 fragment of human AR E6 cDNA (6) was blunt-ended and cloned into the Smal site of pGEX–3X to produce a GST–AR E6 fusion protein. An antiserum against this fusion protein, which cross-reacts to mouse ΔEF1, was used for Western blot analysis.

Immunoprecipitation and Western Blot Analysis. Nuclear extracts were prepared according to the previous report (19). From each 12.5 d.p.c. embryo, 650 μl (2 μg protein/μl) of the extract was obtained, and a 100-μl aliquot was reacted with the anti-AR E6 antisera at room temperature for 2 h. The immunocomplexes were precipitated with protein A-Sepharose (PharMacia) and dissolved in 40 μl of the SDS-PAGE buffer. Each 10-μl sample was separated by SDS-PAGE (7.5% polyacrylamide) and blotted onto a nitrocellulose filter. The blot was treated with 5% skim milk in TBS (20 mM Tris-HCl, 150 mM NaCl, 0.05% Tween 20, pH 8.0), incubated with the anti-AR E6 antisera, washed by TBS, reacted with HRP-conjugated goat anti-rabbit IgG and processed for ECL chemiluminescence reaction (Amersham).

FACS Analysis. Multicolor analysis of lymphocytes was performed by FACScan® cell sorter as described previously (20). The following mAb were purchased from Pharmingen (San Diego, CA): fluorescein isothiocyanate (FITC)-conjugated anti-CD45R, RA3-6B2; FITC-conjugated anti-CD8a, 53-6.7; FITC-conjugated anti-CD3e, 145-2C5; FITC-conjugated anti-Gr-1, R6B6C5; FITC-conjugated anti-CD25, 7D4; PE-conjugated anti-CD4, R M4-5; PE-conjugated anti-α/β TCR, H57-597; PE-conjugated anti-Mac-1, M 1/70; PE-conjugated anti-CD44, 1M 7; Biotin-conjugated anti-LgM, R 6-602, biotin-conjugated anti-κ-kappa receptor, 2B8. Biotin-conjugated anti-α4 integrin (CD49d), M FR 4B. FITC-conjugated anti-IgD was purchased from Nordic Immunological Laboratories (Capistrano Beach, CA). Three-color analysis for κ-kappa, α4 integrin, and CD44/CD25 expression was done using Cy-Chrome-labeled streptavidin (Pharmingen) as the third fluorescent dye and analyzed by FACScan® cell sorter.
Histology and Immunohistology. Tissues were fixed in Bouin’s fixative and paraffin sections were stained with hematoxylin and eosin. For immunofluorescent staining of thymus, a thymus was excised from an 18.5 days post coitus (d.p.c.) embryo, and rinsed in Hepes-buffered saline (HBS), quickly frozen by dipping into liquid nitrogen, and then embedded in OCT compound (Miles Inc., Elkhart, IN). 6-μm-thick cryosections were fixed in 1% paraformaldehyde in HBS for 30 min. The sections were incubated with affinity-purified rabbit anti-ΔEF1 antibodies and biotinylated rat monoclonal anti-CD4 and anti-CD8 antibodies, then with FITC-labeled anti-rabbit Ig and Texas red-conjugated streptavidin anti-rat Ig in TBST containing 10% skim milk with washings by TBS between the steps. Finally, the specimens were mounted in Gelvatol (PBS containing 20% polyvinylalcohol, 20% glycerol, and 2.5% 1,4 diazabyclo-[2,2,2]-octane) and examined under a microscope.

Results

Expression of ΔEF1 in Embryo and in Lymphoid Organs. A cDNA coding for mouse ΔEF1 was isolated by cDNA hybridization with chicken ΔEF1 cDNA that was isolated and characterized in our laboratory (9). Comparison of the mouse ΔEF1 sequence with the chicken and other species revealed that the two zinc finger clusters and the homeodomain are highly conserved (9). Expression pattern of the mouse ΔEF1 in the embryos was almost identical to that of chicken (4): mesodermal tissues (e.g., notochord, somite, limb bud mesenchyme); neural crest derivatives (e.g., dorsal root ganglia, cephalic ganglia); a part of the central nervous system (hindbrain, motor neurons in the spinal cord). In Northern blot analysis of adult tissues, ΔEF1 mRNA was detected in all solid tissues examined (data not shown).

Among the lymphoid tissues, the ΔEF1 transcripts were detected in thymocytes, but not in splenocytes consisting of only mature T and B cells (Fig. 1 A, lanes 2 and 3). Bone marrow cells contained a low but detectable level of the transcripts (Fig. 1 A, lane 1). Immunohistological analysis showed that ΔEF1 protein was expressed in most of the thymocytes including cells stained with the mixtures of anti-CD4 and anti-CD8 antibodies (Fig. 1 B). Taken together with the results of Northern blot analysis, it was indicated that ΔEF1 is expressed in some of the bone marrow cells and in most of the thymocytes including CD4- and CD8-expressing cells, but once the cells migrate out ΔEF1 expression is lost.

Generation of ΔEF1 Mutant Mice Which Lack the Zinc Finger Cluster Proximal to COOH Terminus of ΔEF1 Protein. To generate the ΔEF1 ΔC-fin mutant allele, we constructed the targeting vector shown in Fig. 2 A. It was expected that the product of the recombinant gene lacks the C-proximal zinc finger cluster required for high affinity DNA binding (Fig. 2 B and see reference 12). Germ-line male chimeras were produced from homologous recombinant E14 ES cells (e.g., A84 in Fig. 2 C), and heterozygous mutant animals were generated by crossing these chimeras with C57BL/6 or ICR female mice. The heterozygous mice appeared normal in growth, fertility, behavior, and morphology of internal organs. Southern blot analysis of the yolk sac DNA from the 12.5 d.p.c. embryos generated by crossing the heterozygous mice showed the set of the hybridizing bands expected for the homozygous embryos (Fig. 2 C). Then we examined the ΔEF1 mRNA from the homozygous mutant embryos (12.5 d.p.c.) and compared them with the littermates of other genotypes. As shown in Fig. 2 D, transcripts from the mutated allele were longer than those from wild-type allele as expected from insertion

Figure 1. ΔEF1 expression in adult lymphoid tissues. (A) Total RNAs were prepared from splenocytes (lane 1), thymocytes (lane 2), and bone marrow cells (lane 3) of 8-wk-old wild-type C57BL/6 mice. Each 5-μg RNA sample was analyzed by Northern blotting using a mouse ΔEF1 cDNA as probe. The position of the origin of electrophoresis, ΔEF1 mRNA, and ribosomal RNAs are indicated. The same filter was rehybridized for glyceraldehyde-3-phosphate dehydrogenase (G3PDH) mRNA to control the amount of loaded RNAs. (B) A section of thymus of 18.5 d.p.c. embryo was doubly stained with anti-ΔEF1 antibody (green) and a mixture of anti-CD4 and anti-CD8 antibodies (orange). Note that the majority of the thymocytes had ΔEF1 in the nuclei, together with CD4/CD8 on cell surface. Bar, 10 μm.

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of the neor sequence. The heterozygous mutant embryos had both transcripts while the homozygous mutant embryos possessed only the longer transcript. To further confirm the mutation of δEF1 gene by homologous recombination, we examined the expression of δEF1 protein in the mutant embryos by Western blot analysis (Fig. 2E). The mutant protein is expected to be smaller by 40 kD than the wild-type δEF1 protein. It was demonstrated that wild-type and homozygous mutant embryos had only full-length and truncated forms of δEF1, respectively, while the heterozygous embryos had both. All these observations indicated that δEF1 gene was mutated as designed, and that the truncated form of the mutant protein was synthesized no less efficiently than the wild-type form.

Severe Hypocellularity in the Thymus of δEF1 Mutant Mice. In embryonic stages the homozygous mutant embryos developed up to 18.5 d.p.c. with the expected Mendelian frequency. In histological analysis of the embryos from 10.5 to 18.5 d.p.c., we noted that the thymi of the homozygous mutant embryos were smaller than normal embryos. Other tissues and organs were normal in morphological and histological inspections. The number of thymocytes of 18.5 d.p.c. homozygous mutant embryos was reduced 10-fold (~5 × 10⁵ cells per thymus) compared to the heterozygous and wild-type embryos (~5 × 10⁶ cells per thymus). FACS® analysis of the thymocytes using CD4 and CD8 markers indicated that the development from the DN to DP cells appeared to be partially inhibited (data not shown). We observed no difference between wild-type and homozygous mutant embryos.

In the postnatal period ~80% of the homozygous mutant pups died within 2 d after birth, but the remaining 20% survived and some of them had offspring. After backcrossing to C57BL/6 mice for 6 generations, 11 of the homozygous mutant mice which survived and aged from 3 to 11 wk were analyzed for lymphoid tissue development. As observed in the fetuses, thymi of all the homozygous mutant mice inspected were greatly reduced in size. In histo-

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**Figure 2.** δC-fin δEF1 mutant allele generated by homologous recombination. (A) The last three exons (6–8) of the mouse δEF1 gene encoding the homeodomain and the C-proximal zinc finger cluster are shown (top) together with the targeting vector (middle) and the resulting genomic structure of the homologous recombinant (bottom). Stop codons and the neor cassette are inserted in the middle of the 6th exon, downstream of the homeodomain in the targeting vector. A DT-A cassette (11) was added at the 3’ end of the vector for negative selection against random insertion of the vector. neor, neomycin resistance gene cassette; DT-A, the Diphtheria toxin A chain expression cassette. The diagnostic BglII fragments detected in Southern blots using the probe (indicated by the thick bar) are shown. Restriction sites of SalI, ApaI, and Sau3AI in the genomic DNA used for the targeting vector construction are also shown (see Materials and Methods). (B) Proteins coded by wild-type allele (wt) and the mutated (ΔC-fin) allele are schematically shown. (C) DNAs isolated from wild-type mice (+/+) , a recombinant ES clone (A84), and mice heterozygous (+/−) or homozygous (−/−) for the mutant δEF1 gene were digested with BglII and subjected to Southern blot analysis using the indicated probe. (D) Total RNAs (5 µg each) from wild-type (+/+ ), heterozygous (+/−), and homozygous (−/−) embryos (12.5 d.p.c.) were analyzed by Northern blotting using a mouse δEF1 cDNA (clone M12) as probe. Only the larger size mRNA (δEF1+neo) resulting from the insertion of neo was detected in a homozygous embryo, while only the normal size of δEF1 mRNA was present in a wild-type, and both were in a heterozygous embryo. (E) Nuclear extracts from wild-type (+/+), heterozygous (+/−) and homozygous (−/−) 12.5 d.p.c. embryos were immunoprecipitated and analyzed by Western blotting for δEF1 and ΔC-fin protein using anti-δEF1 antisera which can react to N-proximal portion of δEF1 (see Materials and Methods). As size references, the nuclear extracts from the COS cells transfected with expression vectors of wild-type (wt/COS) and ΔC-fin δEF1 protein (ΔC-fin/COS) were also electrophoresed in parallel.
logical sections of the mutant thymi, medulla and cortex were hardly distinguishable (Fig. 3, A and B); especially, the cortex which usually consists of the densely packed and actively proliferating small thymocytes seemed to be missing in the mutant thymi (Fig. 3, C and D). Accordingly, the total cell number of thymus was greatly reduced from 1/100 to 1/500 of the heterozygous littermate (Fig. 4 A). The spleens of homozygous mutant mice were not significantly different in size from those of heterozygous littermates, although the number of splenocytes was slightly lower in homozygous mutant mice (Fig. 4 C). As will be shown below, the extent of reduction of the cell number seems to be accounted for by the reduction of peripheral T cells in the spleen. Histological inspection of the spleens of the homozygous mutant mice showed the basic architecture to be normal (data not shown). The lymph node of the homozygous mutant mice was characteristic in the reduced cellularity of the deep cortex where the T cells reside (data not shown). The number of lymphocytes recovered from a pair of inguinal lymph nodes of those mice was reduced to ~1/10 of that from the heterozygous littermates (Fig. 4 B).

The heterozygous and wild-type animals were indistinguishable in their histology and cell content of all the lymphoid organs described above. In addition, αEF1 mutant of another allele, Null-LacZ, which lacked almost all of the coding sequence, also exhibited the reduced size of the thymi and impaired T cell development at fetal stages as observed in the ΔC-fin mutant mice (Takagi T., H. Kondoh, and Y. Higashi, unpublished results). These observations exclude the possibility that the T cell defect in the homozygous mutant mice is due to some general cell proliferation defect.

Figure 3. Histology of thymus of αEF1 mutant mouse. Thymi of 6-wk-old heterozygous control (A, C) and homozygous mutant (B, D) mice were fixed in Bouin's fixative and stained with hematoxylin and eosin. The control thymus had developed distinct medulla and cortex (A), while mutant thymi had uniform parenchyma with light staining as seen in medulla of the control thymus (B). Note also the differences of size and cellularity between them. The control thymus had a typical cortex which consists of the densely packed and actively proliferating small thymocytes as shown in higher magnification (C), while the mutant thymus seemed to lack its architecture (D). Bars: (A and B) 200 μm; (C and D) 40 μm.

Figure 4. Total cell count of lymphocytes in lymphoid organs of αEF1 mutant mice in comparison with control heterozygous littermates. Total lymphocyte numbers in thymus (A), spleens (B), and lymph nodes (C) of 6–11 wk were plotted. Note severe reduction of the total lymphocyte numbers in the mutant thymi (~100-fold) and lymph nodes (~10-fold) while the difference in cell number was less pronounced in the spleen.
mozygous ΔC-fin mutant mice is due to the specific effect of the truncated protein.

A aberrant T cell development in the thymus and reduced T cell population in the peripheral lymphoid organs in ΔEF1 mutant mice. Using FACS® with various cell surface markers, we first investigated the cell populations in thymi from the homozygous mutant and heterozygous control mice. In the homozygous mutant mice, the total thymocyte number was so small (see Fig. 4 A) that whole thymocytes of a mutant mouse was subjected to the FACS® analysis. A representative set of the results is shown in Fig. 5 A. The proportion of the CD4/CD8 SP or DP cell populations present was different from that found in a control heterozygous littermate: the relative proportion of cells in CD4⁺CD8⁻ quadrant was reduced from 88% in normal to 57% in the mutant thymocytes while that of the SP cells was increased from 11 to 38% (Fig. 5 A). The relative frequency of α/β TCR⁺CD3⁺ cells in the mutant was higher than in the control heterozygous littermate (Fig. 5 B), which was consistent with the fact that the proportion of SP cells was higher in the mutant mice compared with the control heterozygous littermate as described above.

From forward light scattering data of the FACS® analysis, which indicated the distribution pattern of cell size of a cell population examined, we found that the cell population of larger size predominantly existed in the homozygous mutant mice whereas most of the thymocytes in control mice consisted of the small cell population typical for normal mice (Fig. 5 C). The bias toward the larger size of the cell population in the mutant thymocytes seemed to be due simply to the reduced cell number of the DP thymocytes relative to that of SP cells in the mutant mice since we noticed that the most abundant cell size of the DP and SP cells in the control and mutant mice were not quite different (Fig. 5 C). However, cell size distribution pattern of DN cell population was significantly different between the control and mutant thymocytes: mutant thymocytes were not abundant in the cell population of larger size, but instead had much smaller cells compared to the control mice (Fig. 5 C).

T cells in the spleen and the lymph node were also examined by FACS®. In the mutant spleen, although decrease of total cell number in the mutant was less conspicuous than in the thymus (Fig. 4 B), the fraction of the T cells in the total splenocytes was significantly reduced from ~40% (heterozygous) to 10% (mutant) as judged by Thy1 (T cell marker) and B220 (B cell marker) expression (Fig. 6 A).

In the lymph node, T cells were reduced in number (Fig. 7 A) and were α/β TCR⁺CD3⁺ (Fig. 7 C), indicating full maturity, but the ratio of CD4⁺CD8⁻ SP to CD4⁻CD8⁺ SP cells at the age of 3–8 wk was 5 in the mutant as compared to the ratio of 2 in normal (Fig. 7 B). This bias toward CD4⁺CD8⁻ cell population tended to be greater in

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**Figure 5.** FACS® analysis of thymocytes from a ΔEF1 mutant and a control heterozygous littermate. Thymocytes from 6-wk-old homozygous mutant (left) and heterozygous littermate (right) were analyzed by staining with the combination of mAbs: (A) PE-anti-CD4 vs. FITC-anti-CD8; (B) PE-anti-α/β TCR vs. FITC-anti-CD3 to assess developmental stages of thymocytes. Numbers in parentheses indicate the percentage of cells within the quadrant defined by fluorescence of cell surface markers. The thymocytes were also analyzed by the forward light scattering for the estimation of cell size (C). The histograms for whole or a portion of the thymocytes that were logically gated for the DN, DP, and SP cells in A are shown with combination of heterozygous (+/−) and mutant (−/−) thymocyte data: abscissa, forward light scattering and ordinate, relative cell number.
older mice (e.g., 20 wk, data not shown). In such old mice, the bias also became evident in the spleen though there seemed to be no such bias in the spleen of young adult mice at the age of 3–8 wk (Fig. 6B). The decrease of the T cells commonly observed in the peripheral lymphoid organs may be ascribed to the limited supply of T cells from the thymus.

Since MHC class I and II antigens on the thymic stromal cells are essential for the generation of the CD4 and CD8 SP T cells, we examined the expression of the class I (H-2K) and class II (I-A) antigens on the thymic epithelial cells by immunohistochemistry using anti H-2K and I-A antibodies. The levels of expression of these antigens were not different between ΔEF1 ΔC-fin mutant and control mice, although
The expressions of both MHC antigens were detected in medullary region in the control mice while in the whole region in the mutant mice (data not shown). This difference may suggest and consistent with the histological observation (Fig. 3) that the small thymi in the mutant mice lacked the typical cortex comprising the densely packed small thymocytes, most of which are DP and do not express the MHC antigens highly.

Next we examined the functional maturity of the peripheral T lymphocytes by the proliferative response to the Con A stimulation using the spleen cells as shown in Table 1. Although the degree of \( \Delta \text{cpm} \) of the mutant mice was about one-third of that of the control mice, it seemed that the decrease in \( \Delta \text{cpm} \) in the mutant mice was simply due to reduction of the number of T cells in the spleen cells of homozygous mutant mice (Fig. 6). The sensitivity to Con A stimulation and the size of colonies of the proliferating T cells were not different between the control and mutant mice (data not shown). Thus, functional maturity of the peripheral T lymphocytes accumulated in the \( \Delta \text{EF1} \) homozygous mutant mice did not seem to be affected at least in the responsiveness to the Con A stimulation.

In contrast to T cells, the development of B cells and myeloid cells was not significantly affected by the \( \Delta \text{EF1} \) mutation. B cells were comparable in number among the spleens of homozygous, heterozygous and wild-type littermates. The level of IgM and IgD expression in splenocytes was normal in the homozygous mutant mice (Fig. 6, C and D). The population of the B cells and myeloid cells in the bone marrow was normal as analyzed using B220 (low expression), Mac1 and Gr-1 markers (data not shown). Hematocrit value of the homozygous mutant mice was also similar to the wild-type animals (data not shown). Thus, the defects in the \( \Delta \text{EF1} \) mutant mice appeared specific to T lymphocytes.

Depletion of \( CD4^{-}CD8^{-}c\text{-kit}^{+} \) thymocytes in \( \Delta \text{EF1} \) mutant mice. The FACS\textsuperscript{\textregistered} analysis of the mutant thymocytes using CD4 and CD8 markers indicated that, despite the severe decrease of the total cell number, intrathymic development of the T lymphocytes was not arrested at a specific stage. This is in contrast to the cases of knockout mice lacking molecules that are essential for the development and function of T lymphocytes (e.g., TCR-\( \beta \) [21], CD3 [22]). We suspected that the T precursor cells at a very early stage, for instance, before the rearrangement of the TCR genes, might be affected and decreased in the \( \Delta \text{EF1} \) mutant mice, and that a small fraction of the T precursor cells which escaped from the block by the \( \Delta \text{EF1} \) mutation proceeded to subsequent development.
A nate represents relative cell number. The abscissa indicates intensity of Cy-chrome fluorescence, and ordinate indicates number of cells sampled in each cell population.

To address this point, we analyzed the c-kit expression in the CD4<sup>+</sup>CD8<sup>+</sup> DN cell population by three-color FACS analysis. The early intrathymic T precursor cells which migrate from the bone marrow express c-kit receptor as other hematopoietic progenitors (23, 24). As shown in Fig. 8, A and B, only 15% of the CD4<sup>+</sup>CD8<sup>+</sup> cell population in the mutant thymus was assigned as c-kit<sup>+</sup>, while >50% of the CD4<sup>+</sup>CD8<sup>+</sup> cells were c-kit<sup>+</sup> in the control hematopoietic littermate. Taking into account the difference in the total thymocyte number (Fig. 4) and in the proportion of DN cells (Fig. 5 A), it was concluded that early T cell precursor was depleted in the thymus of δE1 F1 mutant mice.

We analyzed further the DN cell population by staining CD44 and CD25 antigens. It is known that a combination of expression states of these markers defines a stage of thymocyte development within the DN cell population which proceeds in the following order: CD44<sup>+</sup>CD25<sup>+</sup> → CD44<sup>+</sup>CD25<sup>+</sup> → CD44<sup>+</sup>CD25<sup>+</sup> → CD44<sup>+</sup>CD25<sup>+</sup> (25, 26). The CD44<sup>+</sup>CD25<sup>+</sup> population contains the earliest intrathymic T precursor cells, but also include the cells of non-T cell lineages. It is also known that the CD44<sup>+</sup>CD25<sup>+</sup> cells are largest in size and express c-kit most highly, while advancement of the cells to CD44<sup>+</sup>CD25<sup>+</sup> and CD44<sup>+</sup>CD25<sup>+</sup> stages results in smaller cell size and in lower c-kit expression (26, 27). As shown in Fig. 8 C, CD44<sup>+</sup>CD25<sup>+</sup> cells, which correspond to the high c-kit<sup>+</sup> population was greatly reduced from 12 to 3% in the mutant thymocytes as expected. We also noticed that the relative proportion of CD25<sup>+</sup> cells in DN cell population was reduced at least to 50% of that of control mice.

A augmentation of α4 integrin expression in δE1 F1 mutant mice to cause this severe reduction of early T cell precursors? Given that δE1F1 can be a repressor of E2-box sequence (7, 12), genes known to have E2-box sequences in their promoter region are good candidates as a regulatory target of δE1F1. Among them, α4 integrin gene is especially interesting since it contains multiple E2-box sequences (28), and it is known to have an essential role in lymphocyte migration which may include a pathway from bone marrow to thymus (29, 31). Furthermore, α4 integrins have recently been suggested to have some functions in intrathymic development of T lymphocytes (32, 33).

We thus examined the expression of several cell surface markers including integrins in the mutant thymocytes to look for the expression of those proteins affected by δE1 F1 mutation. We found that the expression of α4 integrin (CD49d) was significantly increased (Fig. 9): two- to three-fold increase in α/βTCR<sup>+/low</sup>/CD3<sup>+</sup>/low immature cells of the mutant, and to a lesser extent in α/βTCR<sup>high</sup>/CD3<sup>high</sup> more mature cells (Fig. 9).

Discussion

The data presented in this report demonstrates that δE1 F1 is essential for normal T cell development. In δE1 F1 mutant mice, total lymphocyte number in thymus was reduced to ~1% of that of normal mice. The cell populations found in the thymus of the mutant mice, though reduced in number, represented those of T cells with advanced development, such as CD4<sup>+</sup>CD8<sup>+</sup> DP and CD4<sup>+</sup>CD8<sup>+</sup> SP cells. This phenotype is in contrast to the previously reported mutant mice of T cell development, e.g., RAG (34, 35), TCR<sup>α</sup>β (21, 36), CD3 (22, 37). In these previous cases where the mutated genes were essential for critical steps of T cell development, advancement of thymocytes to subsequent stages was blocked resulting in accumulation of the cells arrested at the critical stages. The total thymocyte number tended to be smaller as the blocked stage became earlier. In the RAG-2-deficient mice, for instance, thymocytes were decreased 100-fold as was observed in δE1 F1 mutant mice, and the existing thymocytes were mostly DN cells (34). The fact that the majority of thymocytes in δE1 F1 mutant mice expressed CD4, CD8 markers despite the severe reduction in total thymocyte number led us to postulate that the majority defect of this mutant lay at a very early stage, much earlier than that at which the TCRs were required, and a small fraction of the cells that somehow escaped from the δE1 F1 mutation could go through the maturation stages.

The idea that the δE1 F1 mutation impairs early T cell development was supported by the observation on c-kit<sup>+</sup> cells in the DN cell population that are considered to be the earliest intrathymic T precursor cells migrating from bone marrow. 

Figure 9. FACS<sup>®</sup> analysis of α4 integrin expression in thymocytes of a δE1 F1 mutant (B) and a heterozygous littermate (A) were stained with FITC-anti-CD3 mAb, PE-anti-CD4 mAb and biotin-conjugated anti-α4 integrin mAb plus Cy-chrome streptavidin, and analyzed for α4 integrin expression in α/βTCR<sup>+/low</sup>/CD3<sup>+</sup>/low α/βTCR<sup>high</sup> CD3 high populations marked by 1 and 2, respectively. Histograms of the α4 integrin expression of mutant (+/−) and heterozygous littermate (+/−) thymocytes were compared in C and D for populations 1 and 2, respectively. Each histogram was drawn so that the total number of cells sampled in each cell population was equal between the mutant and heterozygous littermate. Abscissa indicates intensity of Cy-chrome fluorescence, and ordinate represents relative cell number.
marrow (24). This particular cell population was largely depleted from the mutant thymocytes (Fig. 8 B). Consistent with this observation, the CD4\(^+\)CD25\(^-\) cells, a subset of cells in the early stage of the DN cell population expressing c-kit strongly (26), were also significantly reduced (Fig. 8 C). The cell size distribution of the mutant DN thymocytes measured by the forward light scattering was shifted to smaller than control (Fig. 5 C), which is again in agreement with the observation mentioned above because the c-kit\(^+\) DN thymocytes have been shown to have the largest cell size in the DN cell population (26). Furthermore, δEF1 was expressed in the thymocytes (Fig. 1 A, lane 2) and in the bone marrow cells (Fig. 1 A, lane 1), though at a low level in the latter. All these observations support the idea that δEF1 plays an essential role in the very early stages of T cell development so that abrogation of δEF1 results in the severe decrease of T cell populations accompanied by the significant loss of immature (CD4\(^-\)CD8\(^-\)c-kit\(^+\)) intrathymic T precursor cells.

It is interesting to note that the recently reported mutant mouse of IL-7 receptor (IL-7R) (38), IL-2 receptor γ-chain (IL-2Rγ) (39), and Jak3 (40-42), all showed severe reduction in the thymocyte number, and yet produced all D N, DP, and SP cells, similar to δEF1 mutant mice. Since IL-7R, IL-2Rγ, and Jak3 molecules are thought to be essential signaling molecules in expansion of the thymocytes at a very early stage before rearrangement of the TCR genes, it would be interesting to see if δEF1 is involved in such a signaling pathway.

Besides the early stage of T cell development, δEF1 may have additional regulatory roles in the later stage of intrathymic T cell development. The proportion of D N cells in total thymocytes was higher, and the ratio of D P to SP cells was lower in the δEF1 mutant mice. It was also noted that the mutant thymus did not develop medullo-cortical distinction, probably, due to lack of the typical cortex which usually consists of the densely packed and actively proliferating small thymocytes. The defects of δEF1 mutants are different in these two points from those of IL-7, IL-2Rγ, and Jak3 mutant mice: the proportion of the four cell populations marked by CD4, CD8 expression are identical to the normal mice and the thymi of those mice develop distinct structures of cortex and medulla, though the thymi themselves were very small and the thymocyte numbers are ~1% of the normal mice as the δEF1 mutant mice (38, 39). These differences support the involvement of δEF1 in not only early but also late stages of thymocyte development. R econstitution of the hematopoietic system in wild-type host using mutant hematopoietic tissues will clarify if the defects in early and late T cell development can be ascribed to lymphocytes or stroma.

In peripheral lymphoid tissues the proportion of CD4\(^-\)CD8\(^+\) T lymphocytes relative to CD4\(^+\)CD8\(^-\) cells was significantly reduced from ~0.5 (wild type) to 0.2-0.1 (mutant). In older mice, the majority of the peripheral T cell population was occupied by CD4\(^+\)CD8\(^-\) cells (data not shown). Since the MHC class I and II antigens, which are required for generation of the functional CD4\(^+\)CD8\(^+\) and CD4\(^-\)CD8\(^-\) SP T cells, were expressed at the normal level on thymic epithelial cells of the mutant mice (data not shown), and actually the CD4\(^+\)CD8\(^-\) SP T cells were produced in the mutant thymus (Fig. 5 A), survival of mature CD4\(^+\)CD8\(^+\) cells may be affected in peripheral tissues of δEF1 mutant mice.

An interesting observation was the increase of α4 integrin (CD49d) expression in the mutant thymocyte (Fig. 9). δEF1 may be a truncated form of δEF1-1 integrin, which seems to be a truncated form of δEF1, was reported to function as a repressor of the IL-2 gene transcriptional regulation in human T cell jnk or actin (46). The impairment of T cell development observed in the δEF1-deficient mice could result from the incomplete regulation of the IL-2 gene expression. Deregulated expression of the IL-2 gene in transgenic mouse system, however, did not show any T cell deficiency (47, 48). M oreover, the IL-2 gene knockout mice showed no effects on thymocyte and peripheral T cell subsets (49). T hus, δEF1 mutant phenotype is not explained by altered regulation of the IL-2 gene.

It has been reported that δEF1 (and its human homologue, ZEB) is expressed in myeloma and B-lymphoma cell line (7). T his is not totally consistent with our observation that δEF1 mRNA were not detected in the N orthern blot analysis of the splenocytes. It is possible that expression of δEF1 might be correlated to neoplastic transformation of B cells, or, alternatively, that δEF1 mRNA in mature B or T cells are too little to be detected by our N orthern blot analysis using total R NA.

Although we still do not know the reason for neonatal death observed in part of the δC-fin mutant mice, it was rather unexpected that homozygous mutant embryos carrying the δC-fin allele developed normally except for the T cell defect, given the fact that δEF1 acts as a competitive repressor against bHLH activator proteins, which are widely involved in embryogenesis (12). It should be mentioned that Sna, a mouse homologue of the Drosophila snail and esrigt, is expressed at high levels in cephalic neural crest, limb bud mesenchyme, and somites of 9.5 d.p.c. embryo and later in a variety of mesenchymal tissues (50), showing close resemblance to the expression pattern of δEF1. M oreover, consensus binding sequence of Drosophila snail and...
escargot proteins contain CACCTG E2 box sequence and actually these proteins counteracts E2-box-mediated activation by heterodimer of Suate and D aughters bHLH proteins (51, 52). It is possible that loss of normal $\delta$EF1 protein was compromised by the mouse Sna protein.

In conclusion, T cells, but not other hematopoietic lineage cells, are depleted in $\delta$EF1 mutant mice although they are thought to be derived from common stem cells. Possibly, $\delta$EF1 has an important role in growth and differentiation of early T cells in the bone marrow, in the homing process and/or in development in the thymus. To date, the early T precursor cells have been understood only poorly. $\delta$EF1 mutant mice should thus provide a unique tool to study early T cell development from the aspect of transcriptional regulation.

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