

accelerated aging (39). Furthermore,  $p21^{Waf1/Cip1}$  appears to play protective effects on stem cell exhaustion and ionizing radiation, which may underlie the long-term homeostasis throughout entire life span (40–42). Thus, it is likely that the role of  $p21^{Waf1/Cip1}$  in organismal aging is context-dependent and more complicated than envisaged.

Another important aspect of the  $p53/p21^{Waf1/Cip1}$  pathway is the induction of cellular senescence (4, 24), the state of permanent cell cycle arrest provoked by a variety of oncogenic stimuli including oncogenic *ras* expression in cultured primary cells (43–45). Because this mechanism is poorly understood *in vivo* (46), we tested this notion using our *in vivo* BLI system. Indeed, DMBA/TPA treatment, which causes an oncogenic mutation of the endogenous *H-ras* gene (25, 26), resulted in a significant induction of  $p21^{Waf1/Cip1}$  expression accompanied by skin papilloma formation (Fig. 2A). Because the induction of  $p21^{Waf1/Cip1}$  expression can be seen from the very beginning of papilloma formation and was further enhanced as the papillomas developed (Fig. 2A), the  $p21^{Waf1/Cip1}$  gene is likely to play multiple roles in preventing benign skin papilloma formation and/or malignant conversion. These results are somewhat consistent with two conflicting reports; one shows that disruption of the  $p21^{Waf1/Cip1}$  gene accelerates benign skin papilloma formation but not malignant conversion (27), and the other shows that disruption of the  $p21^{Waf1/Cip1}$  gene accelerates malignant conversion but not benign skin papilloma formation (28).

During the time-course BLI experiments in skin tumor formation, we unexpectedly found that the expression of  $p21^{Waf1/Cip1}$  oscillates over the 3-week time course of the hair follicle cycle (Figs. 2C and D and 3A). Moreover, although the morphology of the four main hair types was not remarkably different, the proportion of zigzag hairs, which are produced by the smallest hair bulb, was significantly reduced in  $p21^{+/-}$  mice (Fig. 3B). Surprisingly, the effects of  $p21^{Waf1/Cip1}$  deficiency were less pronounced in  $p21^{-/-}$  mice than in heterozygotes (Fig. 3B) because of an up-regulation of  $p27^{Kip1}$  expression in the hair bulb of  $p21^{-/-}$  mice (Fig. 3C). In concordance with this notion, recent study revealed that the proportion of zigzag hairs is substantially reduced in mice lacking  $p27^{Kip1}$  (33). Thus, there is cross-talk between  $p21^{Waf1/Cip1}$  and  $p27^{Kip1}$  in controlling hair phenotype. Note that the proportion of zigzag hairs is also reduced in mice lacking the *Sox18* gene (47), and Sox family transcription factors are known to activate  $p21^{Waf1/Cip1}$  expression (48). Thus, it is tempting to speculate that Sox18 controls hair phenotype through regulating  $p21^{Waf1/Cip1}$  gene expression in the hair bulb. It is also important to note, however, that the expression of p63 (34), a member of p53 family transcription factor, overlapped with that of  $p21^{Waf1/Cip1}$  in the hair bulb (Fig. 3A). It is therefore likely that  $p21^{Waf1/Cip1}$  gene expression is controlled by multiple factors in hair bulbs.

We are currently uncertain about the biological consequence of alteration of the zigzag hair proportion in  $p21^{+/-}$  mice. However, because the hair coat plays a crucial role in controlling body temperature in the wild, it is interesting to speculate that  $p21^{Waf1/Cip1}$  and/or  $p27^{Kip1}$  may be involved in temperature control in the wild. Taken together, our results reveal an unexpected role for  $p21^{Waf1/Cip1}$  and provide an insight into how the hair phenotype is determined. Visualizing the dynamics of  $p21^{Waf1/Cip1}$  gene expression in living mice, therefore, provides a powerful tool for not only help to resolve and clarify issues connecting *in vitro* studies but also reveals unrecognized functions of this key proliferative regulator in various physiological processes *in vivo*.

## Materials and Methods

**Generation of p21-p-luc transgenic mouse.** The 2.5-kb fragment of the  $p21^{Waf1/Cip1}$  gene promoter containing two p53-binding sites (GenBank accession number: NW\_923073.1; from 9663447 to

9665888) was placed in front of the firefly luciferase reporter cDNA in the pGL3-basic plasmid vector (Promega, Madison, WI) (see plasmid construction map in SI Fig. 10). The plasmid DNA was digested with XhoI and Sall, and the DNA fragment containing the  $p21^{Waf1/Cip1}$  gene promoter, luciferase cDNA, and polyA signal was isolated and used for microinjection. The transgenic mouse strain was generated by pronuclear microinjection of the reporter transgene into fertilized CD1 oocytes. Genotyping of transgenic mice was determined by PCR and was confirmed by whole-body BLI. One transgenic line, #34 (CDB0415T-34), was selected for studies described in this article because the levels of bioluminescence signals were well correlated with those of endogenous  $p21^{Waf1/Cip1}$  mRNA. All animals were cared for by using protocols approved by the Committee for the Use and Care of Animals of the University of Tokushima.

**Bioluminescence Imaging.** For the detection of luciferase expression, mice were anesthetized, injected i.p. with D-luciferin sodium salt (75 mg/kg) 5 min before beginning photon recording (16–20). Mice were placed in the light-tight chamber, and a gray-scale image of the mice was first recorded with dimmed light, followed by acquisition of luminescence image by using a cooled CCD camera (Princeton Instruments, Trenton, NJ). The signal-to-noise ratio was increased by  $2 \times 2$  binning and 5 min exposure, unless otherwise stated in the text. For colocalization of the luminescent photon emission on the animal body, gray scale and pseudocolor images were merged by using IMAGE-PRO PLUS (Media Cybernetics, Bethesda, MD).

**Tumor-Induction Experiments.** Twelve mice of the  $p21$ -p-luc line, in the resting phase of the hair cycle (8-week-old), were shaved and treated with 7,12-dimethylbenzanthracene (DMBA) (100  $\mu$ g in 100  $\mu$ l of acetone). One week after DMBA treatment, mice were subsequently treated twice a week with TPA (12.5  $\mu$ g in 100  $\mu$ l of acetone) for 20 weeks (25, 26). Control mice were treated with acetone instead of DMBA/TPA.

**Semiquantitative RT-PCR.** Total RNA was isolated by using TRIzol reagent (Invitrogen, Carlsbad, CA), and 2  $\mu$ g of total RNA was used for the reverse-transcriptase reaction. The PCR was performed by using Blend Taq polymerase (TOYOBO, Osaka, Japan) with primers specific for the mouse  $p21^{Waf1/Cip1}$  gene and the mouse  $\beta$ -actin gene. The PCR primer sequences used are shown in SI Text.

**Real-Time RT-PCR.** Quantitative real-time RT-PCR was performed by using the SYBER Premix EX Taq system (TAKARA, Otsu, Japan) and an ABI Prism 7900HT (Applied Biosystems, Foster City, CA). Amplified signals were confirmed to be single bands by gel electrophoresis and were normalized to the levels of GAPDH. Data were analyzed by using SDS2.1 software (Applied Biosystems). The PCR primer sequences used are shown in SI Text.

**Histology and Immunohistochemistry.** Biopsies of mouse skin taken from the middorsal region were fixed in 10% formalin for a 24 h or longer, progressively dehydrated through gradients of alcohol, and embedded in paraffin. Samples were sectioned on a microtome (5- $\mu$ m-thick), deparaffinized in xylene, rehydrated, and then stained with hematoxylin and eosin (H&E). The hair cycle status was determined by histological examination. For antibody staining, deparaffinized and rehydrated sections were exposed to heat-induced antigen retrieval for 5 min or 20 min in 10 mM citrate buffer (pH 6.0). After washing in PBS, endogenous peroxidase activity was quenched for 15 min in 1%  $H_2O_2$  in methanol, followed by washing with PBS. The sections were



incubated in blocking serum for 1 h at room temperature. After incubation with primary antibodies overnight at 4°C, biotinylated anti-mouse secondary antibody was applied and detected by the avidin-biotin peroxidase technique using the DAB kit (DAKO, Glostrup, Denmark) and then counterstained with methyl green. For immunofluorescence, the relevant Alexa Fluor 488 goat anti-mouse or 546 goat anti-rabbit antibodies (1:1,000; Invitrogen) were used for detection of primary antibodies. Fluorescence images were observed and photographed by using an immunofluorescence microscope (Carl Zeiss, Oberkochen, Germany). The primary antibodies used are shown in *SI Text*.

**Hair Measurements.** Hair was removed from the middorsal region of each mouse (8-week-old) to reduce the potential regional variation, although this is not known to occur in the dorsal region of the mouse. One hundred hairs were examined to determine the percentage of each hair type.

We thank Dr. P. Leder (Department of Genetics, Harvard Medical School, Boston, MA) for providing *p21<sup>Waf1/Cip1</sup>* knockout mice. We also thank Ms. S. Chiba for her assistance in various mouse experiments. This work was supported by grants from Ministry of Education, Science, Sports, and Culture of Japan, the Astellas Foundation for Research on Metabolic Disorders and the Astellas Foundation for Research on Medical Resources.

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## Crosstalk between RANKL and Fas signaling in dendritic cells controls immune tolerance

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Although receptor activator of nuclear factor (NF)- $\kappa$ B ligand (RANKL) signaling has been shown to prolong the survival of mature dendritic cells (DCs), the association of RANKL pathway with Fas-mediated apoptosis is obscure. Here, we found that bone marrow-derived DCs (BMDCs) from the Fas-deficient strain *MRL/lpr* mice, could survive much longer than normal DCs. The expressions of Bcl-x and Bcl-2 and the nuclear transport

of NF- $\kappa$ B of RANKL-stimulated BMDCs from *MRL/lpr* mice were significantly up-regulated. By contrast, Fas expression of BMDCs from normal C57BL/6 and *MRL<sup>+/+</sup>* mice was increased by RANKL stimulation, and an enhanced DC apoptosis was found when stimulated with both RANKL and anti-Fas mAb, which was associated with activation of caspase-3 and caspase-9. Furthermore, the expression of FLIP<sub>L</sub>, an inhibitory molecule against Fas-mediated

apoptosis, in normal DCs was significantly decreased by RANKL and anti-Fas mAb. Indeed, the adoptive transfer of RANKL-stimulated DCs resulted in rapid acceleration of autoimmunity in *MRL/lpr* recipients. These findings indicate that the crosstalk between RANKL and Fas signaling in DCs might control immune tolerance. (Blood. 2007;110:242-250)

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### Introduction

Dendritic cells (DCs) are professional antigen-presenting cells that reside in peripheral tissues in an immature state for optimal antigen uptake and respond to inflammatory signals.<sup>1,2</sup> After capture of pathogen-derived antigens and activation by pathogens, DCs migrate from tissues to the lymphoid organs where DCs become mature and initiate antigen-specific T cells.<sup>1,2</sup> While many subpopulations of DCs have been reported to be heterogeneous in their phenotype and localized in lymphoid tissues, there is general agreement that DCs originate from a hematopoietic progenitor, and that there are 3 DC subtypes, including myeloid DCs (CD11c<sup>+</sup>CD11b<sup>+</sup>CD8 $\alpha$ <sup>-</sup>), lymphoid DCs (CD11c<sup>+</sup>CD11b<sup>-</sup>CD8 $\alpha$ <sup>+</sup>), and plasmacytoid DCs (CD11c<sup>+</sup>B220<sup>+</sup>Gr-1<sup>+</sup>), all of which have been recently identified.<sup>3-6</sup>

The lifespan of activated antigen-bearing DCs has been estimated to be as short as 3 days.<sup>7,8</sup> Limiting the lifespan of DCs by means of apoptosis may serve to regulate the availability of antigen for T cells and to control immune responses. An important and well-characterized mechanism for apoptosis in immune cells is mediated by Fas, the death receptor.<sup>9</sup> Although Fas has been implicated in the apoptosis of DCs,<sup>10,11</sup> DCs are known to be resistant to Fas-induced cell death.<sup>12</sup> In addition, it has been reported that the resistance to Fas-induced apoptosis in DCs correlates with the constitutive expression of the Fas-associated death domain-like IL-1 $\beta$ -converting enzyme (FLICE)-inhibitory protein (FLIP) ligand.<sup>13</sup> However, the molecular mechanism for maintenance of DCs through Fas-mediated signals is still unclear.

Receptor activator of nuclear factor (NF)- $\kappa$ B ligand (RANKL),<sup>14,15</sup> a type II membrane protein of the tumor necrosis factor (TNF) family, is expressed on osteoblasts, stromal cells, and

activated T cells,<sup>15</sup> and binds to the signaling receptor RANK and the decoy receptor osteoprotegerin (OPG).<sup>15,16</sup> Mice lacking RANKL or RANK display dramatically reduced osteoclastogenesis, show defects in early differentiation of T and B cells, systemic lymph nodes, and fail to develop mammary glands.<sup>17,18</sup> Moreover, recent data have showed that the interaction of Fas ligand and Fas expressed on osteoclast precursors increases RANKL-induced osteoclastogenesis.<sup>19</sup> On the other hand, RANKL-RANK interaction has been shown to prolong the survival of mature DCs, whereas the association of the RANKL pathway with Fas-mediated signals in the activation or function of DCs has been obscure.<sup>15,20</sup>

In this study, to define the molecular mechanism for the autoimmunity by immune dysregulation of DCs through Fas and the RANKL pathway, the *MRL/lpr* mouse strain, an autoimmune-prone strain that has a mutated Fas gene, was used to analyze the immune functions of the DCs and autoimmunity such as rheumatoid lesions.

### Materials and methods

#### Mice

*MRL/Mp-lpr/lpr* (*MRL/lpr*; aged 4-12 weeks; n = 105), *MRL<sup>+/+</sup>* mice (aged 4-12 weeks; n = 55), and C57BL/6 (B6; n = 50) mice were purchased from Charles River Japan Inc. (Atsugi, Japan). All mice were maintained in specific pathogen-free conditions in our animal facility, and the experiments were approved by an animal ethics board of Tokushima University (Tokushima, Japan).

Submitted November 28, 2006; accepted March 8, 2007. Prepublished online as Blood First Edition paper, March 19, 2007; DOI 10.1182/blood-2006-11-059980.

The online version of this article contains a data supplement.

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### Generation of murine BMDCs

A crude population of DCs was generated *in vitro* from mouse bone marrow as described with some modifications.<sup>21,22</sup> Briefly, bone marrow was flushed from the long bones of the limbs. Cells were plated at a density of  $1 \times 10^6$  and were cultured for 7 days in 5% CO<sub>2</sub> at 37°C in RPMI 1640 medium containing 10% heat-inactivated fetal bovine serum (FBS), 100 U/mL penicillin, 10 ng/mL recombinant mouse granulocyte-monocyte colony-stimulating factor (GM-CSF; PeproTech, Le Parray-en-Yvelines, France), and 5 ng/mL mouse IL-4 (PeproTech). DCs (CD11c<sup>+</sup> cells, < 70%) including mainly immature DCs (50% CD11c<sup>+</sup>CD86<sup>-</sup> cells, < 50%) were routinely used for the experiments.

### Flow cytometric analysis

Bone marrow-derived DCs (BMDCs) were stained with fluorescein isothiocyanate (FITC)-conjugated anti-CD11c mAb, along with rat anti-major histocompatibility complex class II (MHC class II; I-A<sup>k</sup>) as primary Ab and phycoerythrin (PE)-conjugated anti-rat IgG as secondary Ab. BMDCs were also stained with PE-anti-CD86, PE-anti-CD80, or anti-Bcl-2, anti-Bcl-xL, and PE-anti-rat IgG. For each staining, isotype-matched mAb was used as a control. All mAbs were obtained from BD Biosciences (San Diego, CA). The cells were analyzed with an EPICS flow cytometer (Beckman Coulter Inc, Miami, FL), and data were analyzed with FlowJo FACS analysis software (Tree Star Inc, Ashland, OR).

### Detection of apoptotic cells

Cells were treated with or without 100 ng/mL Fas-activating antibody (Jo-2; BD Biosciences) for 24 to 72 hours and were detected with an EPICS flow cytometer using an Annexin V-FITC apoptosis detection kit (Genzyme, Cambridge, MA). Briefly, after cultured cells were washed in phosphate-buffered saline (PBS), the cells were incubated with FITC-conjugated Annexin-V and propidium iodide (PI) for 10 minutes at room temperature in the dark. Binding buffer was added, and apoptotic cells were detected by flow cytometric analysis with an EPICS flow cytometer.

### Measurement of cytokine production

Cytokine production was tested by a 2-step sandwich enzyme-linked immunosorbent assay (ELISA) using a mouse IL-12p40, IL-10, and interferon (IFN)- $\gamma$ , IL-2, and IL-4 kit (Genzyme, Cambridge, MA). In brief, the culture supernatants from DCs or T cells were added to microtiter plates precoated with anti-IL-12p40, IL-10, IFN- $\gamma$ , IL-2, and IL-4 capture Ab and incubated overnight at 4°C. After adding biotinylated detecting Ab and incubating at room temperature for 45 minutes, streptavidin-peroxidase was added and incubated again at room temperature for an additional 30 minutes. Finally, an 2,2'-azino-di-3-ethylbenzthiazoline sulfonate substrate containing H<sub>2</sub>O<sub>2</sub> was added, and the colorimetric reaction was read at an absorbance of 450 nm using an automatic microplate reader (Bio-Rad Laboratories Inc, Hercules, CA). The concentration was calculated according to the standard curves produced by various concentrations of recombinant cytokines.

### MTT assay

The effect of RANKL on proliferation of BMDCs was determined by the MTT dye uptake method. Briefly, the cells (2000/well) were incubated in triplicate in a 96-well plate in the presence or absence of RANKL in a final volume of 0.1 mL for the indicated time periods at 37°C. Then, 0.025 mL MTT solution (5  $\mu$ g/mL in PBS) was added to each well. After incubation at 37°C for 2 hours, 0.1 mL of the extraction buffer (20% sodium dodecyl sulfide [SDS], 50% dimethylformamide) was added. Incubation was continued overnight at 37°C, and then the optical density (OD) was measured at 450 nm using an automatic microplate reader (Flow, McLean, VA), with the extraction buffer as blank.

### Western blot analysis

The cell extracts from the nucleus and cytoplasm of DCs were prepared using the Nuclear/Cytosol Fractionation Kit (Bio Vision, Mountain View,

CA). Cells were briefly washed, collected in ice-cold PBS in the presence of phosphatase inhibitors, and centrifuged at 230g for 5 minutes. The pellets were resuspended in a hypotonic buffer, treated with detergent, and centrifuged at 14 000g for 30 seconds. After collection of the cytoplasmic fraction, the nuclei were lysed and nuclear proteins were solubilized in lysis buffer containing protease inhibitors. A total of 50  $\mu$ g of each sample per well was used for SDS-polyacrylamide gel electrophoresis. After blocking with 5% nonfat milk, the membrane was incubated with primary antibodies for TRAF6, IKK, p-I $\kappa$ B $\alpha$ , NF- $\kappa$ B p65 (RelA), and p50 (Santa Cruz Biotechnology, Santa Cruz, CA). Antibody-antigen complexes were detected using a horseradish peroxidase-conjugated secondary antibody. Protein binding was visualized with enhanced chemiluminescence Western blotting reagent (Amersham Biosciences, Arlington Heights, IL). Western blotting using anti-Bcl-2, anti-Bcl-xL, anti-Bax, anti-Bid (BD Bioscience), anti-caspase-3, anti-caspase-8, and anti-caspase-9 Abs (Cell Signaling Technology, Beverly, MA), and anti-FLIP<sub>L</sub> (Santa Cruz Biotechnology) was performed essentially as described for the lysis buffer. Control for protein loading was provided by anti-mouse histone, or GAPDH mAb (Sigma Chemical Co, St Louis, MO).

### siRNA of Fas and RANK

For small interfering RNA (siRNA) of Fas and RANK, a siTrio Full Set (B-Bridge International, Sunnyvale, CA) was used for analyzing the function of DCs. Briefly, each cocktail including the 3 RNA oligonucleotides listed here was transfected into cells with a Silencer siRNA Transfection II kit (Ambion, Austin, TX). Sequences of the oligonucleotide sets are as follows: Fas, GGGAAAGGAGUACAUGGACATT (sense), UGUCCAUGUACUCCUCCCTT (antisense), CGAAAGUACCGGA-AAAGAATT (sense), UUCUUUCCGGUACUUUCGTT (antisense), CCAGAAGGACCUUGGAAAATT (sense), UUUUCCAAGGUCCUUCUGGTT (antisense); and RANK, CCAAGGAGGCCAGGCUUATT (sense), UAAGCCUGGGCCUCCUUGGTT (antisense), GGGAAAG-CGUGACAGCUATT (sense), UAGCUGUCAGCGCUUCCCTT (antisense), CUGAAAAGCACCUGACAAAATT (sense), UUUGUCAG-GUCUUUUCAGTT. Transfected cells were incubated with or without RANKL, and flow cytometric analysis was performed.

### Confocal microscopic analysis

Immature DCs were seeded onto glass-bottom culture dishes (MatTek, Ashland, MA) at a density of 500 cells/well and stimulated for 30, 60, or 120 minutes with RANKL. Subsequently, cells were washed and fixed in cold 3% paraformaldehyde (PFA) in PBS for 10 minutes, then permeabilized with 0.2% Triton-X in PBS for 2 minutes. The slides were then washed thoroughly with PBS and incubated with optimal dilutions of the primary Ab in PBS containing 1% bovine serum albumin (BSA)-2.5% FBS in PBS for 1 hour at room temperature. Cells were stained with optimal dilutions of the primary Abs for 1 hour. After 3 washes with 0.0001% Triton-X in PBS, the cells were stained with Alexa Fluor 488 goat anti-rabbit IgG (H + L; Molecular Probes, Eugene, OR) or Texas red-conjugated goat anti-mouse (Molecular Probes) as the second Abs for 30 minutes and washed with PBS. For each fluorochrome label, negative control Abs were added. Finally, the slides were incubated with 4', 6-diamidino-2-phenylindole (DAPI) for 5 minutes to label the nuclei and mounted in Vectashield (Vector Labs, Burlingame, CA) for analysis by confocal microscopy (LSM5 PASCAL, Carl Zeiss, Gottingen, Germany). A 63  $\times$  1.4 oil DIC objective lens was used. Quick Operation Version 3.2 (Carl Zeiss) for imaging acquisition and Adobe Photoshop CS2 (Adobe Systems, San Jose, CA) for image processing were used.

### Transfer of DCs

BMDCs were stimulated *in vitro* for 48 hours with 100 ng/mL RANKL and 50  $\mu$ g/mL bovine type II collagen (CII). RANKL-stimulated BMDCs were transferred into the base of the tail by subcutaneous injections in 200  $\mu$ L PBS at the age of 4 weeks.



## Histopathology

All organs were removed from the mice, fixed with 4% phosphate-buffered formaldehyde (pH 7.2), and prepared for histologic examination. The sections (4  $\mu$ m in thickness) were stained with hematoxylin and eosin (HE). Histologic grading of inflammatory arthritis was performed according to the methods by Edwards et al<sup>23</sup> as follows: a 1-point score indicates hyperplasia/hypertrophy of synovial cells, fibrosis/fibroplasia, proliferation of cartilage and bone, destruction of cartilage and bone, and mononuclear cell infiltrate. Imaging was analyzed by microscope (BX50, Olympus, Tokyo, Japan) at 20  $\times$  10.70 objective lens. Viewfinder Lite Version 1.0 (Olympus) for image acquisition and Adobe Photoshop CS2 for image processing were used.

## Measurement of anti-dsDNA Ab, RF, and CII Ab levels

Anti-double-stranded DNA (dsDNA) Abs, rheumatoid factor (RF), and anti-CII Ab were detected by ELISA as described previously.<sup>11</sup> Briefly, flat-bottom plates (Nalge Nunc International, Roskilde, Denmark) were coated with 1.5  $\mu$ g/mL of native calf thymus DNA (Life Technologies, Rockville, MD) in buffer containing 0.1 M sodium bicarbonate and 0.05 M citric acid at 4°C overnight. Serum samples were serially diluted (starting at 1:200) and added to the plates for 1 hour of incubation at 37°C. After washing, peroxidase-conjugated goat anti-mouse IgG, or IgM (Southern Biotechnology Associates, Birmingham, AL) was added and incubated for 1 hour at 37°C. Ab binding was visualized using orthophenylenediamine (Sigma Chemical). For the measurement of IgG and IgM RF, human IgG and IgM (Chemicon International, Temecula, CA) were coated onto plates at 10  $\mu$ g/mL (pH 9.6). The microtiter plate was coated with 100  $\mu$ L CII antigen solution. After washing 3 times, 100  $\mu$ L/well of serum samples that had been serially diluted in PBS/Tween 20/1% BSA and control serum samples were added and incubated for 1 hour at 37°C. After washing, peroxidase-conjugated goat anti-mouse IgG (at 1.4  $\mu$ g/mL, 100  $\mu$ L/well; Organon Teknika, Durham, NC) was added and incubated for 1 hour at 37°C. A total of 100  $\mu$ L *o*-phenylenediamine (0.5 mg/mL) dissolved in 0.1 M citrate buffer (pH 5.0) containing 0.012% H<sub>2</sub>O<sub>2</sub> was added, and the reaction was stopped using 8 N H<sub>2</sub>SO<sub>4</sub> (20  $\mu$ L/well).

## Statistics

The Student *t* test was used for statistical analysis. *P* values greater than .05 were considered significant.

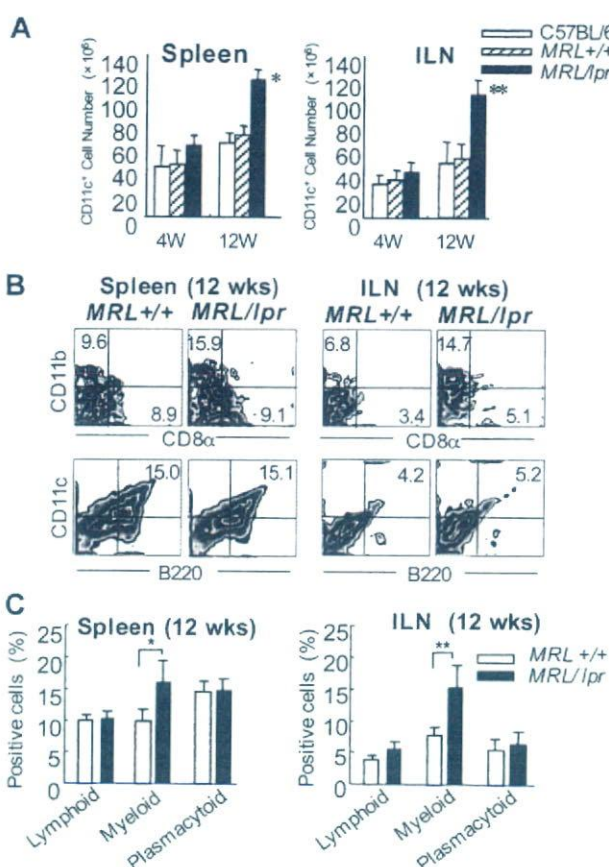
## Results

### DC numbers and subtypes of *MRL/lpr* mice

To examine the number of DCs from *MRL/lpr* mice, the spleen cells and inguinal lymph node (ILN) cells were analyzed at 4 and 12 weeks of age. We found that the DC number of both spleen and ILNs from *MRL/lpr* mice was significantly higher than that from control mice at 12 weeks of age (Figure 1A). We next analyzed the increased subset of DCs from *MRL/lpr* mice. A significant increase of the myeloid (CD11b<sup>+</sup>CD11c<sup>+</sup>CD8 $\alpha$ <sup>-</sup>) DC subset of both spleen and ILNs from *MRL/lpr* mice was observed in contrast to the population from control *MRL*<sup>+/+</sup> mice, whereas there was no difference in the lymphoid (CD11c<sup>+</sup>CD11b<sup>-</sup>CD8 $\alpha$ <sup>+</sup>) and plasmacytoid (B220<sup>+</sup>CD11c<sup>+</sup>) DCs between *MRL/lpr* and control mice (Figure 1B-C). Based on these findings, we recognized the possibility that myeloid DCs from *MRL/lpr* mice may have any influence on the immune disorder of the mice.

### Activation of RANKL-stimulated DCs from *MRL/lpr* mice

To determine the function of DCs from *MRL/lpr* mice, we generated BMDCs by using the culture of bone marrow cells



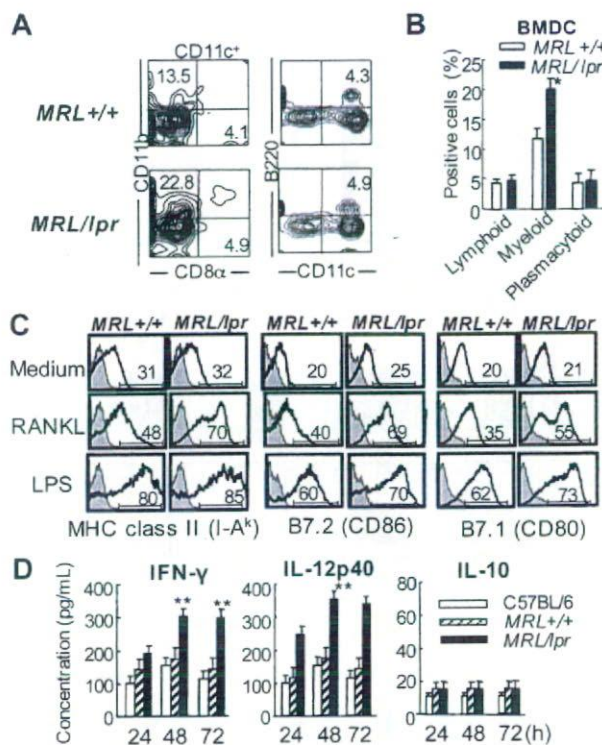
**Figure 1. Myeloid dendritic cells in the spleen and inguinal lymph nodes from *MRL/lpr* mice.** (A) The CD11c<sup>+</sup> DC cell number of the spleen and ILNs is shown as means  $\pm$  SD of 5 to 7 *MRL/lpr* and control mice at 4 and 12 weeks of age. (B) Mononuclear cells were isolated from the spleen and ILNs at 12 weeks of age, and the proportions were determined using flow cytometry. Shown are representative plots of CD8 $\alpha$ <sup>+</sup>, CD11b<sup>+</sup>, B220<sup>+</sup>, and CD11c<sup>+</sup> cells. Percentages in each region indicate the frequency of lymphoid (CD11c<sup>+</sup>CD11b<sup>-</sup>CD8 $\alpha$ <sup>+</sup>), myeloid (CD11c<sup>+</sup>CD11b<sup>+</sup>CD8 $\alpha$ <sup>-</sup>), and plasmacytoid (CD11c<sup>+</sup>B220<sup>+</sup>) DCs. Data are representative of 5 to 7 mice in each group. (C) Graph shows the mean frequency of lymphoid, myeloid, and plasmacytoid DCs. \**P* > .05; \*\**P* > .01 *MRL/lpr* versus *MRL*<sup>+/+</sup> mice. Results are representative of 3 independent experiments.

with recombinant mouse GM-CSF and IL-4. When the cell-surface markers on BMDCs after the culture for 7 days were determined, the myeloid subset DCs from *MRL/lpr* mice had increased significantly compared with that from *MRL*<sup>+/+</sup> mice, resembling the phenotype of the spleen and ILN from *MRL/lpr* mice (Figure 2A-B).

Next, activation of the BMDCs from *MRL*<sup>+/+</sup> and *MRL/lpr* mice stimulated with RANKL or lipopolysaccharide (LPS) was estimated by analyzing the surface phenotypes, including MHC class II, B7.2 (CD86) and B7.1 (CD80). BMDCs were stimulated with 100 ng/mL mouse recombinant RANKL or 100 ng/mL LPS for 48 hours, and then the surface expressions were detected by flow cytometric analysis as shown in Figure 2C. The expressions of MHC class II, B7.1, and B7.2 on BMDCs from *MRL/lpr* mice in response to RANKL were enhanced compared with those from *MRL*<sup>+/+</sup> mice, but no significant difference in the expressions on the BMDCs stimulated with LPS from between *MRL*<sup>+/+</sup> and *MRL/lpr* mice (Figure 2C).

This finding indicates that activation of BMDCs from *MRL/lpr* mice via RANKL might be enhanced. In addition, to define the secretion of some cytokines such as IFN- $\gamma$ , IL-12p40, and IL-10 from RANKL-stimulated BMDCs of *MRL/lpr* mice, BMDCs from





**Figure 2. Surface phenotypes and cytokine productions in RANKL-stimulated DCs from *MRL/lpr* mice.** (A) BMDCs were generated from *MRL/lpr* and *MRL<sup>+/+</sup>* mice by using the culture of bone marrow cells with GM-CSF and IL-4 for 7 days; the cell-surface markers of BMDCs were then analyzed. BMDCs from *MRL/lpr* and *MRL<sup>+/+</sup>* mice were analyzed by flow cytometry. Shown are representative plots of CD8 $\alpha$ <sup>+</sup>, CD11b<sup>+</sup>, B220<sup>+</sup>, and CD11c<sup>+</sup> cells. Percentages in each region indicate the frequency of lymphoid, myeloid, and plasmacytoid DCs. Results are representative of 3 independent experiments. (B) The graph shows the mean frequency of lymphoid, myeloid, and plasmacytoid DCs. Data are means  $\pm$  SD of 5 to 7 mice in each group. (C) BMDCs were stimulated with 100 ng/mL mouse recombinant RANKL or 100 ng/mL LPS for 48 hours, and then the surface expressions of MHC class II and costimulatory molecules were detected by flow cytometric analysis. The result is representative of 3 independent experiments. (D) The secretions of IFN- $\gamma$ , IL-12p40, and IL-10 in culture supernatants from the BMDCs stimulated by RANKL were detected by ELISA. Data are means  $\pm$  SD of triplicate samples and are representative of 4 independent experiments. \* $P > .05$ ; \*\* $P > .01$ , *MRL/lpr* versus control mice.

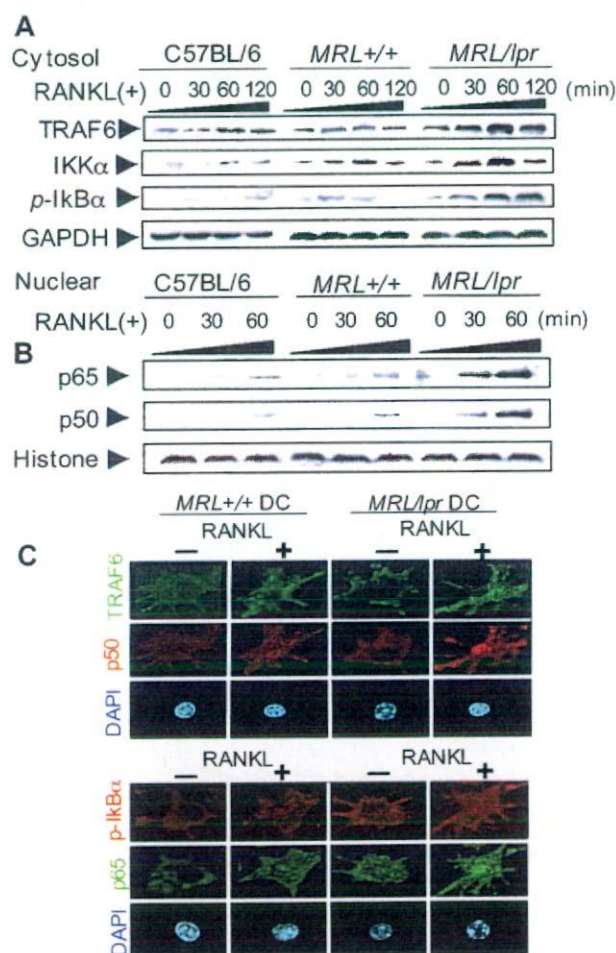
normal B6 mice, *MRL<sup>+/+</sup>*, and *MRL/lpr* mice were stimulated with RANKL for 24 to 72 hours, and the cytokine secretion of the culture supernatants was analyzed by ELISA. Significantly increased secretions of IFN- $\gamma$  and IL-12p40, not IL-10, of BMDCs from *MRL/lpr* mice were observed compared with those from C57BL/6 (B6) and *MRL<sup>+/+</sup>* mice (Figure 2D). These data suggest that the Fas molecule may influence the activation and function of DCs through RANKL signaling.

#### Signaling molecules of DCs through RANKL

It is well known that the RANK/RANKL pathway includes a lot of signaling molecules.<sup>24,25</sup> Among them, NF- $\kappa$ B plays a key role for DC maturation and activation.<sup>26</sup> NF- $\kappa$ B activation through RANK/RANKL occurs by nuclear translocation following inducible phosphorylation of inhibitory I $\kappa$ B by IKK complex.<sup>27,28</sup> Therefore, IKK $\alpha$  expression and phosphorylation of I $\kappa$ B (p-I $\kappa$ B) in RANKL-stimulated BMDCs from B6, *MRL<sup>+/+</sup>*, and *MRL/lpr* mice were analyzed by Western blotting. In addition, the expression of TRAF6 downstream from RANK in RANKL-stimulated BMDCs from *MRL/lpr* and control mice was determined.

Increasing expression of TRAF6 after stimulation with RANKL was detected in BMDCs from *MRL<sup>+/+</sup>* mice. In *MRL/lpr* mice, a

much higher expression of TRAF6 in the BMDCs was observed in contrast to that of the control mice. In addition, the expression of IKK $\alpha$  of *MRL/lpr* DCs was up-regulated relative to those of B6 and *MRL<sup>+/+</sup>* DCs (Figure 3A). Notably, the prominent phosphorylation of I $\kappa$ B of *MRL/lpr* DCs was detected compared with that of control DCs parallel with the increased expressions of TRAF6 and IKK $\alpha$  (Figure 3A). Furthermore, nuclear translocation of NF- $\kappa$ B subunits (p65 and p50) of RANKL-stimulated *MRL/lpr* DCs had increased remarkably compared with that of control DCs (Figure 3B). To define the movements of RANKL-mediated signaling molecules of *MRL/lpr* DCs, the expression of each protein was confirmed by confocal microscopic analysis. Consistent with the results of Western blot analysis using the cytoplasmic and nuclear proteins, the increased nuclear translocation of NF- $\kappa$ B and the up-regulated TRAF6 and p-I $\kappa$ B in the cytoplasm from RANKL-stimulated



**Figure 3. RANKL signaling molecules in *MRL/lpr* DCs.** (A) The expressions of TRAF6, IKK $\alpha$ , and phospho-I $\kappa$ B (p-I $\kappa$ B) of RANKL-stimulated BMDCs from C57BL/6, *MRL<sup>+/+</sup>*, and *MRL/lpr* mice were detected using the cytoplasmic extracts by Western blot analysis. The BMDCs were stimulated with RANKL (100 ng/mL) from 0 to 120 minutes. GAPDH was used as a housekeeping protein. (B) Nuclear translocation of NF- $\kappa$ B (p65 and p50) of RANKL-stimulated BMDCs from C57BL/6, *MRL<sup>+/+</sup>*, and *MRL/lpr* mice was detected by immunoblot. Histone was used as a housekeeping protein of the nuclear extracts. (C) Increased TRAF6 and p-I $\kappa$ B in the cytoplasm, and accelerated nuclear translocation of p50 and p65 in the nucleus of RANKL-stimulated *MRL/lpr* DCs were detected by confocal microscopic analysis. BMDCs from *MRL/lpr* and *MRL<sup>+/+</sup>* mice were stimulated for 48 hours with or without RANKL, fixed in 3% PFA on a glass slide, and stained with anti-p65, anti-p50, TRAF6, and p-I $\kappa$ B $\alpha$  followed by Alexa Fluor 488-labeled (green) or Alexa Fluor 568-labeled (red) anti-mouse or anti-rabbit IgG as the second antibodies. The nuclei were stained with DAPI. Original magnification,  $\times$  630. All data are representative of 3 to 5 independent experiments.



*MRL/lpr* DCs were observed in contrast to those from control DCs (Figure 3C). These findings suggest that the Fas molecule might regulate NF- $\kappa$ B activation of DCs through RANKL signaling.

#### Survival signaling of *MRL/lpr* DCs

To determine the growth of RANKL-stimulated BMDCs from *MRL/lpr* mice, the cell growth was analyzed by MTT assay. At 48 and 72 hours after RANKL stimulation, significantly increased growth of *MRL/lpr* DCs was observed compared with those of B6 and *MRL<sup>+/+</sup>* DCs (Figure 4A).

We next analyzed survival signaling molecules, Bcl-xL and Bcl-2, that are antiapoptotic molecules among the Bcl-2 family by flow cytometry. Enhanced expressions of both intracellular Bcl-xL and Bcl-2 in RANKL-stimulated *MRL/lpr* DCs were observed at 24 and 72 hours after the stimulation in contrast to that in *MRL<sup>+/+</sup>* DCs (Figure 4B). Moreover, prominent amounts of both Bcl-xL and Bcl-2 in RANKL-stimulated *MRL/lpr* DCs were detected compared with those in *MRL<sup>+/+</sup>* DCs using Western blot analysis (Figure 4C). By contrast, the expressions of Bax and Bid, apoptosis signaling molecules, in RANKL-stimulated *MRL/lpr* DCs were lower than those of *MRL<sup>+/+</sup>* DCs (Figure 4C). These data suggest

that RANKL signal leads to enhancing the survival of Fas-deficient DCs through up-regulation of antiapoptotic molecules.

#### RANKL signaling and Fas-mediated apoptosis of DCs

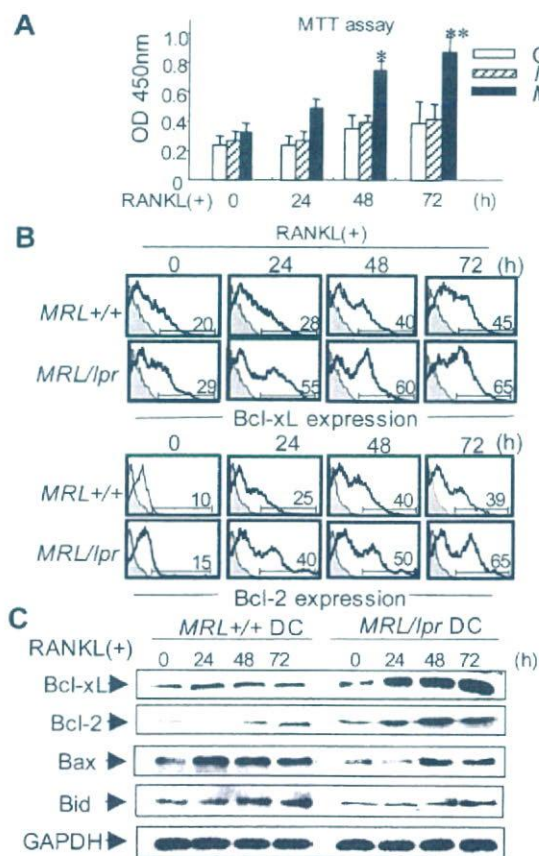
To examine the association of RANKL signaling with Fas-mediated apoptosis in DCs, Fas expression on BMDCs stimulated with RANKL was analyzed. A time-dependent increase in Fas expression on RANKL-stimulated BMDCs from normal B6 and *MRL<sup>+/+</sup>* mice, but not *MRL/lpr* mice, was observed (Figure 5A). We found an enhanced apoptosis of DCs when stimulated with both anti-Fas mAb and RANKL, whereas the apoptosis induced by RANKL or anti-Fas mAb was slightly increased (Figure 5B-C). Moreover, to clarify the molecular pathway of enhanced Fas-mediated apoptosis of DCs by RANKL, the movement of caspases downstream from Fas to undergo apoptosis was analyzed using *MRL<sup>+/+</sup>* DCs by Western blotting. Although a small amount of cleaved caspase-3 and caspase-9 of RANKL-stimulated *MRL<sup>+/+</sup>* DCs was observed, we detected an increased active form of caspase-3 and caspase-9 in the *MRL<sup>+/+</sup>* DCs stimulated with both anti-Fas and RANKL (Figure 5D). The activation of these caspases was clearly inhibited by the addition of caspase inhibitor (z-VAD; Figure 5D). Furthermore, the expression of FLIP<sub>L</sub>, an inhibitory molecule against Fas-mediated apoptosis,<sup>29,30</sup> was significantly decreased by both RANKL and anti-Fas mAb parallel with the movement of the caspases (Figure 5D). It has been reported that FLIP<sub>L</sub> is constitutively expressed in DCs to play an inhibitory role for Fas-mediated apoptosis of DCs.<sup>12,13,31</sup> In addition, the expression of pro-caspase-8 was decreased by the stimulation of RANKL or anti-Fas mAb, and the expression was even lower with stimulation by both RANKL and anti-Fas mAb (Figure 5D). For *MRL/lpr* DCs, the expression of FLIP<sub>L</sub> was not reduced with RANKL stimulation (Figure 5E). These results strongly suggest that RANKL signaling may contribute to Fas-mediated apoptosis of DCs by controlling the expression of FLIP<sub>L</sub>.

#### Effects of Fas siRNA on the function of DCs

To further confirm the enhanced antiapoptotic signal of RANKL-stimulated DCs by Fas deficiency in the *MRL/lpr* mice, we used siRNA of Fas to evaluate the apoptotic signaling and activation of normal BMDCs from control mice.

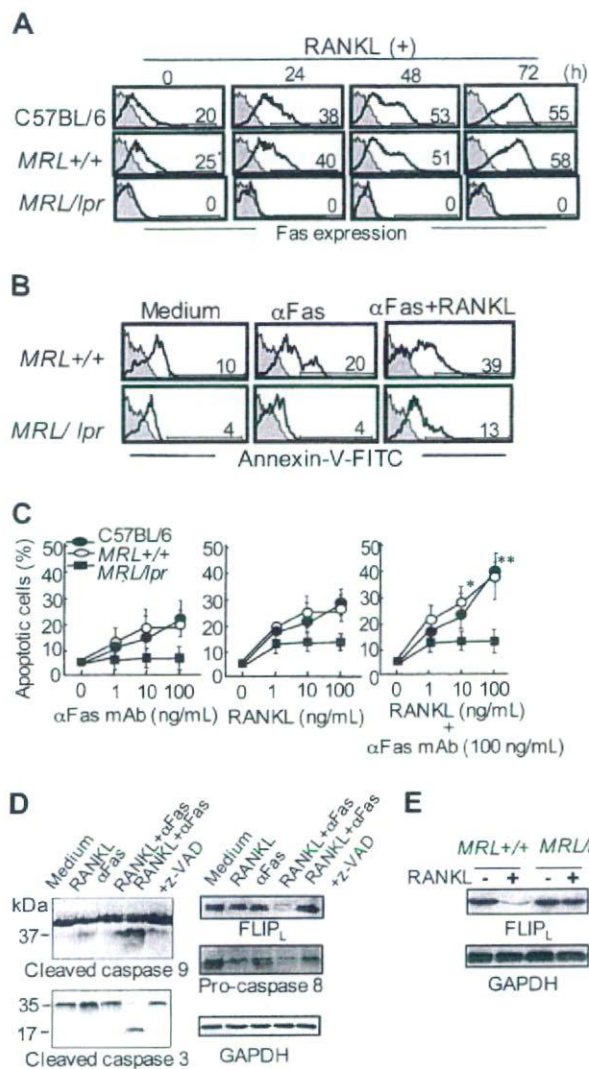
When BMDCs transfected with the siRNA of Fas were stimulated with RANKL for 48 hours, RANKL-induced Fas expression was decreased to the unstimulated level (Figure 6A). We next examined the inhibitory effect of Fas siRNA on RANKL/Fas-induced apoptosis of BMDCs. As expected, apoptosis of RANKL- and anti-Fas mAb-stimulated BMDCs was clearly inhibited by Fas siRNA, but not GAPDH siRNA or an irrelevant control (Figure 6B). Moreover, to clarify apoptosis signal of RANKL/Fas-mediated apoptosis of DCs, cleavage of caspase-3 was examined using Fas siRNA. A cleaved caspase-3 of BMDCs transfected with control siRNA was detected at 48 and 72 hours after RANKL stimulation, but no active form of caspase-3 of BMDCs transfected with Fas siRNA was observed (Figure 6C).

In addition, to confirm the role of Fas signal for the RANKL-induced activation of DCs, MHC class II expression, as 1 of some activation markers on RANKL-stimulated BMDCs treated with the siRNA, was analyzed by flow cytometry. RANKL-induced expression of MHC class II was still more enhanced by the transfection with Fas siRNA, but not control siRNA (Figure 6D). IL-12 secretion from RANKL-stimulated BMDCs of *MRL<sup>+/+</sup>* mice was



**Figure 4. Antiapoptotic signaling through RANKL in *MRL/lpr* DCs.** (A) BMDCs were stimulated with 100 ng/mL mouse recombinant RANKL from 0 to 72 hours, and the cell growth was analyzed by MTT assay. Data are means  $\pm$  SD of triplicate samples and are representative of 4 independent experiments. \* $P > .05$ ; \*\* $P > .01$ , *MRL/lpr* versus control mice. (B) BMDCs from *MRL/lpr* and control mice were stimulated with RANKL (100 ng/mL) from 0 to 72 hours, and the intracellular expressions of Bcl-xL and Bcl-2 were analyzed by flow cytometry. Numbers above horizontal lines indicate percent Bcl-xL<sup>+</sup> or Bcl-2<sup>+</sup> cells. Results are representative of 3 independent experiments. (C) BMDCs from *MRL<sup>+/+</sup>* and *MRL/lpr* mice were treated with RANKL (100 ng/mL) from 0 to 72 hours. Bcl-xL, Bcl-2, Bax, and Bid proteins were detected by Western blot analysis. GAPDH was used as a control for loading. Data are representative of 3 independent experiments.



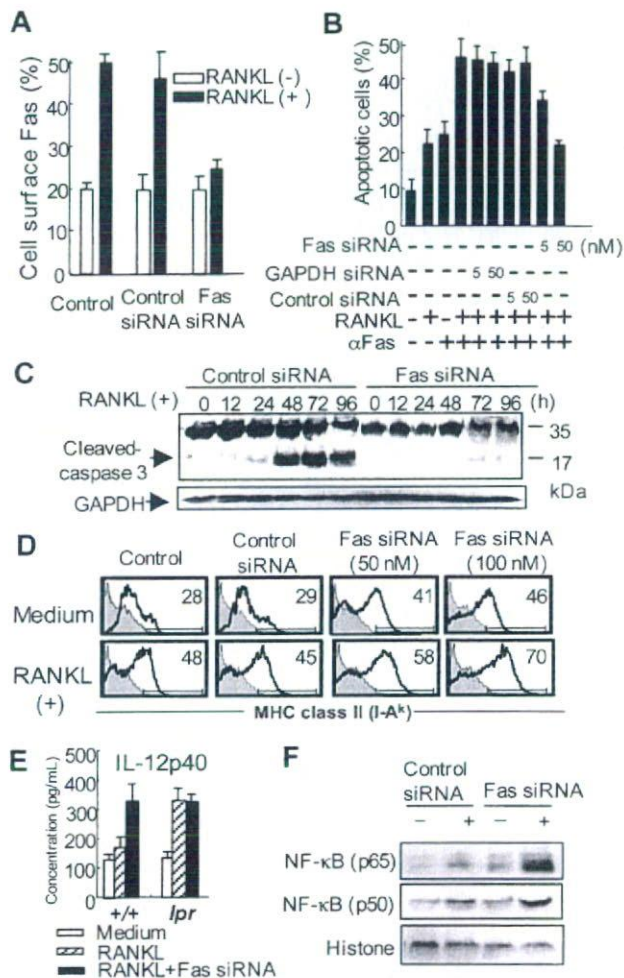


**Figure 5. Fas signaling through RANKL in DCs.** (A) BMDCs from normal B6, *MRL+/+*, and *MRL/lpr* mice were cultured with RANKL from 0 to 72 hours. Fas expression on the DCs was detected by flow cytometry. Numbers above horizontal lines indicate percentage of Fas<sup>+</sup> cells. Results are representative of 3 independent experiments. (B) BMDCs from *MRL+/+* and *MRL/lpr* mice were incubated with anti-Fas mAb, or anti-Fas mAb and RANKL, for 48 hours. Apoptotic cells were detected by flow cytometric analysis. Numbers above horizontal lines indicate percentage of Annexin-V<sup>+</sup> cells. Results are representative of 3 independent experiments. (C) Apoptotic cells stimulated with anti-Fas mAb (1-100 ng/mL), RANKL (1-100 ng/mL), or anti-Fas mAb and RANKL, for 48 hours. B6 (●), *MRL+/+* (○), and *MRL/lpr* (■) mice are shown. Data are means ± SD of 3 independent experiments. \**P* > .05; \*\**P* > .01. (D) BMDCs (5 × 10<sup>4</sup> cells) were cultured treated with RANKL (100 ng/mL) and anti-Fas mAb (100 ng/mL) in the presence or absence of inhibitor z-VAD for 48 hours. The expressions of caspase-9, caspase-3, caspase-8, and FLIP<sub>L</sub> of the stimulated BMDCs from control mice were analyzed by Western blot. GAPDH was used as a control for loading. Results are representative of 3 independent experiments. (E) FLIP<sub>L</sub> expression of RANKL-stimulated BMDCs from *MRL+/+* and *MRL/lpr* mice was detected by immunoblot. GAPDH was used as a control for loading. Results are representative of 3 independent experiments.

elevated by the treatment with Fas siRNA, indicating that Fas deficiency positively controls RANKL signaling as observed in the DCs from *MRL/lpr* mice (Figure 6E). Moreover, the accelerated nuclear translocation of NF-κB in *MRL/lpr* DCs was observed in RANKL-stimulated BMDCs treated with Fas siRNA (Figure 6F). These findings indicate that RANKL-stimulated DCs from normal mice may be maintained by Fas-mediated apoptosis through caspase and NF-κB cascade.

**Inhibitory effect of RANK siRNA on Fas-mediated apoptosis of DCs**

To clarify the direct association between RANK/RANKL and Fas signaling of DCs, Fas-mediated apoptosis of RANK-deficient DCs was analyzed using the siRNA of RANK as shown in the protocol (Figure S1A, available on the *Blood* website; see the Supplemental Figures link at the top of the online article). The knockdown of RANK by the siRNA in BMDCs was confirmed by analyzing the protein expression of RANK on flow cytometry and Western blotting as shown in Figure S1B-C. Then, Fas expression and



**Figure 6. Effects of Fas siRNA on the functions of DCs.** (A) Fas expression on Fas siRNA-transfected BMDCs from control mice was analyzed after the stimulation of RANKL. GAPDH siRNA or irrelevant oligonucleotides was used as control. Data are means ± SD of triplicate samples and are representative of 3 independent experiments. (B) Inhibitory effect of Fas siRNA on Fas and RANKL-induced apoptosis of BMDCs from control mice was observed. Apoptotic cells were detected by flow cytometric analysis. Data are means ± SD and representative of 3 independent experiments. (C) Fas siRNA-transfected BMDCs were stimulated with RANKL from 0 to 96 hours. Caspase-3 expression was analyzed by immunoblot. GAPDH was used as a control for loading. Data are representative of 3 independent experiments. (D) MHC class II expression on the Fas siRNA-transfected BMDCs stimulated with RANKL was detected by flow cytometric analysis. Numbers above horizontal lines indicate percentage of MHC class II<sup>+</sup> cells. Data are representative of 3 independent experiments. (E) Fas siRNA-transfected BMDCs from *MRL+/+* and *MRL/lpr* mice were stimulated with RANKL, and the secretion of IL-12 from the DCs was detected by ELISA. Data are means ± SD of triplicate samples and are representative of 3 independent experiments. (F) Nuclear translocation of NF-κB subunits (p65 and p50) was detected using the nuclear extracts of Fas siRNA-transfected BMDCs from control mice by Western blot analysis. Histone was used as a control for loading. Data are representative of 3 independent experiments.



apoptosis of RANK-deficient DCs from *MRL<sup>+/+</sup>* and *MRL/lpr* mice was analyzed as shown in Figure S1D-E. Expectedly, RANKL-induced Fas expression was decreased to the level of unstimulation in RANK-deficient DCs from *MRL<sup>+/+</sup>* mice (Figure S1D). Furthermore, the Fas- and RANKL-induced apoptosis was reduced to the level of anti-Fas stimulation (Figure S1E).

### Influence of activated *MRL/lpr* DCs on arthritis

It is still unclear whether the activated and increased DCs from *MRL/lpr* mice influence autoimmune lesions such as arthritis in the mice. To determine whether the activated DCs from *MRL/lpr* mice may influence autoimmune lesions in vivo, the adoptive transfer of BMDCs stimulated with RANKL and CII into *MRL/lpr* mice at 4 weeks was performed to analyze arthritis lesions of the recipient mice. The histologic score of arthritis in the mice transferred with RANKL- and CII-stimulated *MRL/lpr* DCs was significantly higher at 12 and 16 weeks of age than those of the mice transferred with nonstimulated *MRL/lpr* DCs, RANKL- and CII-stimulated *MRL<sup>+/+</sup>* DCs, or the untreated mice (Figure 7A). Histologic findings of the recipient mice transferred with RANKL-

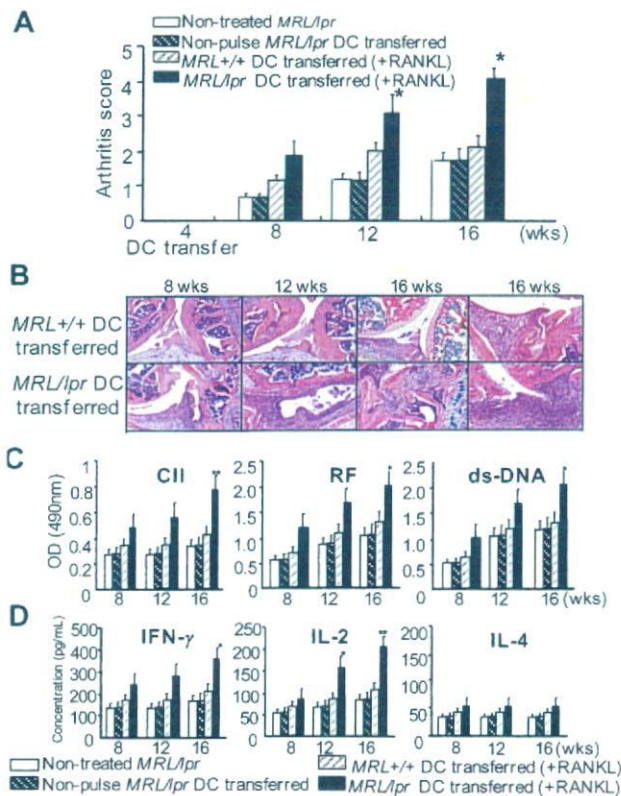
CII-stimulated *MRL/lpr* DCs showed severe subsynovial inflammation, synovial hyperplasia, pannus formation, cartilage erosion, and bone destruction (Figure 7B). In contrast, the histology of the mice transferred with RANKL- and CII-stimulated *MRL<sup>+/+</sup>* DCs was mild, similar to untreated *MRL/lpr* mice (Figure 7B). Anti-CII Ab and RF of sera from the recipient mice transferred with RANKL- and CII-stimulated *MRL/lpr* DCs were significantly increased (Figure 7C). Moreover, increased secretions of IFN- $\gamma$  and IL-2, not IL-4 from T cells stimulated with anti-CD3 mAb of the recipient mice transferred with RANKL- and CII-stimulated *MRL/lpr* DCs, were detected compared with those of the control recipients (Figure 7D). These results indicate that the RANKL-stimulated DCs might play an important role in the acceleration of autoimmune arthritis in *MRL/lpr* mice through T helper 1 (Th1) response.

### Discussion

Studies in the past several years have established an important role of DCs in controlling many immune responses.<sup>1,2</sup> However, the mechanism of maintenance of peripheral DCs has been unclear. The present study indicates that the maintenance of DCs, including cell growth, activation, and apoptosis in the periphery, is regulated by a crosstalk between RANKL and Fas signaling, and that the RANKL/Fas system in DCs may play a key role for the maintenance of peripheral tolerance.

DCs are heterogeneous in terms of phenotype, localization, and function. Several DC subsets have been described based on surface expressions, such as CD11c, B220, CD11b, CD4, and CD8 $\alpha$ .<sup>3,4</sup> It has been reported that plasmacytoid DCs induce Th2 cell differentiation in response to certain stimuli, and that myeloid DCs favor a Th1 response.<sup>6</sup> Among myeloid DCs, 4 subsets are known: CD11b<sup>+</sup>CD8 $\alpha$ <sup>+</sup>, CD11b<sup>+</sup>CD4<sup>+</sup>, CD11b<sup>+</sup>CD4<sup>-</sup>, and CD11b<sup>low</sup>CD8 $\alpha$ <sup>-</sup>CD4<sup>-</sup>.<sup>3,5,6</sup> The function of each subset has been obscure. In this study, increased CD11c<sup>+</sup>CD11b<sup>+</sup>CD8 $\alpha$ <sup>+</sup> myeloid DCs from *MRL/lpr* mice were observed, implying that the maintenance of Th1 DCs in the periphery may occur through Fas signaling, and that Th1 DCs from *MRL/lpr* mice might influence immune disorders such as autoimmune lesions and lymphadenopathy in the mice. It has been described that defective T-cell function of *MRL/lpr* mice plays a key role for pathogenesis of the immune disorder.<sup>10,32,33</sup> Our data demonstrate that dysfunction of DCs from *MRL/lpr* mice accelerates autoimmunity mediated by T and B cells.

It has been reported that the Fas-mediated apoptosis of DCs is resistant due to a constitutive FLIP that can block Fas-induced apoptosis, whereas Fas is expressed on the surface of DCs.<sup>13,31</sup> Upon Fas activation, a set of effector molecules is recruited to the receptor leading to form a signaling complex. Initially, FADD binds to Fas, recruits caspases, and activates a cascade of apoptosis. The present study indicates that Fas-mediated apoptosis of DCs can be induced by the addition of activated RANK/RANKL signaling, suggesting that there may be a crosstalk between the Fas and RANK pathway for DC maintenance in the periphery. In addition, our results indicate that the expression of FLIP<sub>L</sub> of DCs is controlled by RANKL and Fas signaling, leading to NF- $\kappa$ B activation. This is consistent with the recent report that cFLIP controls NF- $\kappa$ B activation and maintenance in lymphocytes and DCs.<sup>34</sup> The molecular mechanism may be mediated by activation-induced cell death (AICD), which is well known to act as a system to maintain the peripheral T cells through Fas-mediated apoptosis.<sup>35-37</sup> Activated or autoreactive T cells are considered to delete by



**Figure 7. Effects of the RANKL-stimulated *MRL/lpr* DCs transfer on autoimmune arthritis.** BMDCs pulsed with CII and stimulated with RANKL were transferred into *MRL/lpr* recipients (4 weeks of age). The recipient *MRL/lpr* mice were analyzed at 8 to 16 weeks of age. (A) Histologic score of autoimmune arthritis in *MRL/lpr* mice transferred with RANKL-stimulated *MRL/lpr* DCs was evaluated compared with control mice until 16 weeks of age. Data are means  $\pm$  SD of 5 to 7 mice in each group. \* $P > .05$ ; \*\* $P > .01$ , *MRL<sup>+/+</sup>* DCs versus *MRL/lpr* DCs transferred. (B) Histologic analysis of autoimmune lesions from *MRL/lpr* mice transferred with RANKL-stimulated *MRL/lpr* DCs and *MRL<sup>+/+</sup>* DCs at 8, 12, and 16 weeks of age was performed. The sections of joints from the recipients were stained with HE. Data are representative of 5 to 7 mice in each group. (C) RF, anti-CII antibodies, and anti-dsDNA antibodies of sera from the recipients at 16 weeks of age were detected by ELISA. (D) The secretions of IFN- $\gamma$ , IL-2, and IL-4 in the supernatants from ILN T cells stimulated with anti-CD3 mAb were detected by ELISA. Data are means  $\pm$  SD of triplicate samples and are representative of 5 to 7 mice in each group. \* $P > .05$ ; \*\* $P > .01$ , *MRL<sup>+/+</sup>* DCs versus *MRL/lpr* DCs transferred.



using AICD. Our data suggest that "AICD" of DCs may be mediated by the Fas and RANK pathway.

Bcl-2 family proteins, including Bcl-2 and Bcl-xL, are considered to be critical for DC survival, and to be regulated by NF- $\kappa$ B.<sup>7,38</sup> In this study, Bcl-xL and Bcl-2 expressions of *MRL/lpr* DCs were considerably up-regulated by RANKL stimulation, leading to hyperproliferation of DCs. In addition, NF- $\kappa$ B activation of *MRL/lpr* DCs induced by RANKL was accelerated, suggesting that the Fas signaling pathway might influence NF- $\kappa$ B activation via RANKL signaling in DCs. The NF- $\kappa$ B family has emerged as a key transducer of inflammatory signals to DC maturation and activation, and is well known to be activated by various stimuli such as RANK/RANKL, CD40, and toll-like receptors (TLRs).<sup>14,39,40</sup> Accelerated signaling pathways, including IKK $\alpha$ , I $\kappa$ B, and NF- $\kappa$ B downstream from TRAF6, was observed in *MRL/lpr* DCs stimulated by RANKL. Therefore, RANKL signaling leading to NF- $\kappa$ B activation through TRAF6 in DCs may be regulated by Fas-induced signaling.

DCs are crucial for the pathogenesis of autoimmune disease because of their potent antigen-presenting activity and unique ability to activate naive T cells.<sup>41</sup> DCs have also been described to prime autoreactive T cells and induce the local inflammation of the synovial membrane in arthritis.<sup>42</sup> It was reported that the transfer of collagen-pulsed BMDCs into congenic DBA/1 mice results in inflammatory arthritis.<sup>43</sup> In addition, antigen-pulsed DCs have been shown to induce disease in experimental autoimmune encephalomyelitis, a murine model of multiple sclerosis.<sup>44</sup> Our study showed that transfer of RANKL-stimulated *MRL/lpr* DCs pulsed with collagen into *MRL/lpr* mice clearly accelerates arthritis in the recipients, suggesting that the activated DCs stimulated by RANKL may accelerate autoreactivity of T and B cells to enhance the productions of pathogenic cytokines and autoantibodies in *MRL/lpr* mice. As for autoimmune arthritis of *MRL/lpr* mice, the increased DCs, especially myeloid DCs in the periphery, may play the key role for the pathogenesis. It was previously reported that an

imbalance favoring development of DCs from myeloid-committed progenitors predisposes to autoimmune lesion in nonobese diabetic (NOD) mice.<sup>45</sup> Therefore, analyzing the differentiation and function of myeloid DCs may lead to understanding the pathogenic mechanism of autoimmune disease.

Taken together, the new finding in the present study is that the maintenance of DCs, including cell growth, activation, and apoptosis in the periphery, is regulated by crosstalk between RANKL and Fas signaling, and that the RANKL/Fas signaling in DCs may play a crucial role in the maintenance of immune tolerance.

## Acknowledgments

The authors thank Ai Nagaoka, Hiroyo Amo, and Satoko Yoshida for technical assistance.

This work was supported by the Ministry of Education, Science, and Culture of Japan (Grants-in-Aid for Scientific Research nos. 17109016 and 17689049).

## Authorship

Contribution: T.I. performed all experiments and wrote the manuscript; N.I. codesigned experiments, coproduced confocal microscopic analysis, and wrote the manuscript; K.M. codesigned experiments and wrote the manuscript; M.K. coproduced ELISA with T.I.; R.A. performed Western blot analysis; Y.H. unified the project and wrote the manuscript.

Conflict-of-interest disclosure: The authors declare no competing financial interests.

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## 特集I

## T細胞レセプターからのシグナル伝達

T細胞レセプター  
シグナルとNF- $\kappa$ B\*石丸直澄\*\*  
岸本英博\*\*\*  
林良夫\*\*Key Words: NF- $\kappa$ B, T cell receptor, I $\kappa$ B, activation, NF- $\kappa$ B2(p100/p52)調節機構が明らかにされ, 自己免疫疾患の病態発症におけるNF- $\kappa$ Bシグナルの謎が解き明かされようとしている.

## はじめに

T細胞の活性化や増殖にはT細胞そのものが産生するサイトカインや増殖因子が必須である. そのサイトカインや増殖因子の転写を制御している転写因子の中でもっとも重要な位置に存在しているのがnuclear factor(NF)- $\kappa$ Bである. NF- $\kappa$ Bは, IL-2をはじめとしたサイトカインおよびサイトカインレセプター, 増殖因子, 細胞接着分子, 細胞周期に関連する遺伝子, アポトーシス関連遺伝子などT細胞の生死にかかわる重要な遺伝子の転写調節を司っている<sup>1)</sup>. 一般的にT細胞受容体(T cell receptor; TCR), CD28を中心とした副刺激分子およびサイトカインレセプターなどを介した刺激からさまざまなシグナル分子を経由してNF- $\kappa$ Bの内在性阻害因子であるI $\kappa$ Bのリン酸化, ユビキチン化および断片化によってI $\kappa$ Bから解離したNF- $\kappa$ B分子のヘテロおよびホモダイマーが細胞質から核内に移行し, 標的遺伝子上に存在する $\kappa$ Bサイトを介して転写が調節され, さまざまな重要因子が合成されT細胞の機能獲得に用いられている. 最近, I $\kappa$ BによるNF- $\kappa$ Bの制御機構に加えて新たにNF- $\kappa$ B2による

2つのNF- $\kappa$ B経路

転写因子であるNF- $\kappa$ Bは免疫細胞における多くの炎症反応を制御している. NF- $\kappa$ BにはN末にDNA結合に重要な共通のRelホモロジー領域を有するNF- $\kappa$ B1(p105-p50), NF- $\kappa$ B2(p100-p52), RelA(p65), RelB, c-Relの5つのサブユニットがあり(図1), それらのサブユニットがヘテロダイマーあるいはホモダイマーとなり核内移行し, 細胞の活性化や増殖などに重要な遺伝子の転写調節を司ることが知られている<sup>2)</sup>. NF- $\kappa$ B経路にはNF- $\kappa$ B1とRelAの複合体によって調節される古典的経路(classical NF- $\kappa$ B pathway)と, NF- $\kappa$ B2とRelBの複合体によって調節される非古典的経路(non-classical NF- $\kappa$ B pathway)が報告されている<sup>3)</sup>(図2).

TCR/CD3関連シグナルとNF- $\kappa$ B

T細胞では主にTCRからの刺激がさまざまな伝達分子を介して古典的NF- $\kappa$ Bの活性化につながるがよく知られている<sup>4)</sup>. まず, TCRからの抗原刺激によりTCR/CD3分子複合体に連動してZAP70, Fyn, Vavやphosphoinositide 3-kinase

\* The critical NF- $\kappa$ B pathway through T cell receptor.

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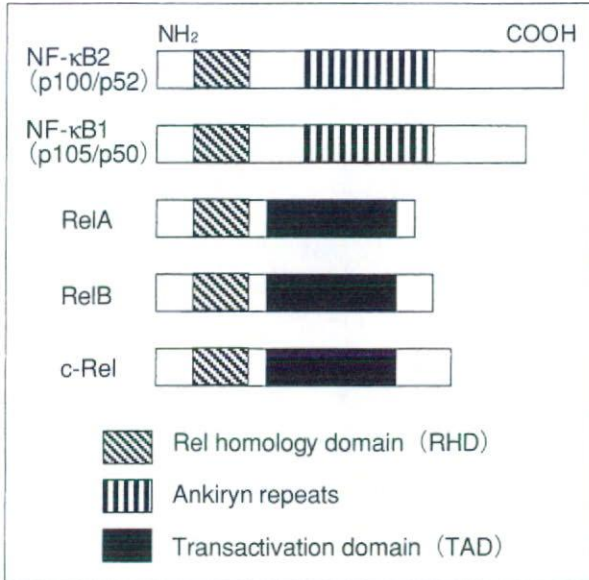


図1 NF-κBサブユニットの構造

(PI3K)といったアダプター分子を介して, protein kinase C θ(PKCθ), caspase-recruitment domain-containing membrane-associated guanylate kinase protein 1(CARMA1)のリン酸化へと進んでいく. PKCθより上流のシグナルはAP-1, NFATといったNF-κBとは別のルートによるT細胞の

活性化経路と共通しているが, 3-phosphoinositide-dependent protein kinase 1(PDK1)から引き続いて誘導されるPKCθとCARMA1のリン酸化はNF-κBの活性化に重要な起点となっている. PKCθのリン酸化に続いてCARMA1, B-cell lymphoma 10(Bcl-10), mucosa-associated-lymphoid-tissue lymphoma-translocation gene 1(MALT-1)複合体を介してIκB kinase (IKK)の活性化につながっていく<sup>5)6)</sup>. また, PKCθが直接IKKに結合してIKKの活性を調節していることも明らかにされている<sup>7)</sup>. 一方, 副刺激分子であるCD28からのシグナルに関しては, TCRの刺激がない状態でCD28刺激を入れてもNF-κBの活性化は不十分であり, T細胞の活性化および増殖といった正常な機能獲得には至らない. CD28分子からのシグナルはTCRの下流のPI3KやAKTなどのアダプター分子を経由して上記のPKC/CARMA1/Bcl-10/MALT-1を介してIKKの活性化につながり, TCRからのNF-κBシグナルを量的に調節しているものと考えられる. IL-2レセプターなどのサイトカインレセプターからのシグナルについてもTCRの下流のシグナル分子と合流していることが知られている<sup>8)9)</sup>.

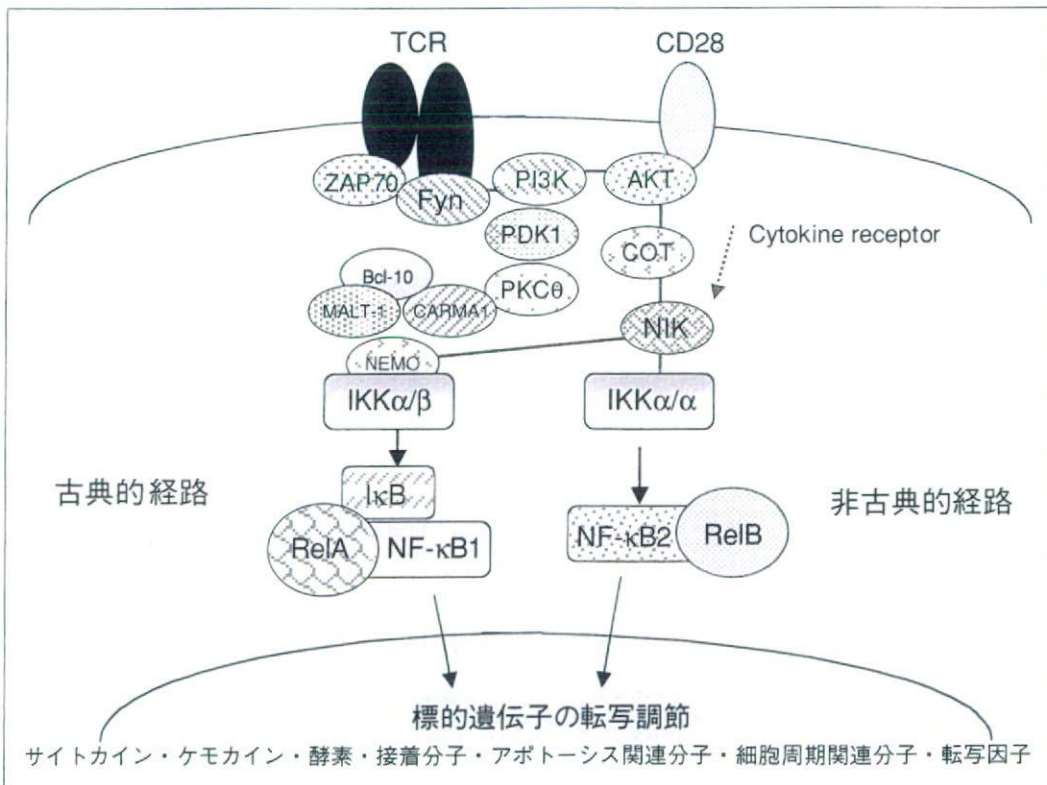


図2 T細胞レセプターと古典的NF-κB経路および非古典的(新規)NF-κB経路 T細胞あるいはCD28の下流におけるNF-κBの転写活性に至る過程の概略図を示す.



### IκBによる制御システム

NF-κBの活性化の引き金となるのが、NF-κB分子と結合しているIκBのリン酸化から始まるユビキチン化、断片化である。通常はNF-κB分子は細胞質でIκBと結合することによって核内に移行しないように制御されている<sup>5)</sup>。CARMA1/Bcl-10/MALT-1複合体の活性化からIKKのキナーゼ活性が高まりIκBのリン酸化が誘導されると、IκB分子はユビキチン化を受けた後プロテアソームにより断片化されることによりNF-κBの核内移行を阻害できなくなる。つまり、IκBによる制御が解除されるときにNF-κBの活性化は開始される。正常マウスのCD4陽性T細胞を抗TCR抗体および抗CD28抗体により刺激すると、IκBのリン酸化および断片化、さらに、NF-κB分子の核内移行が観察される(図3)。IκB分子をリン酸化するのがIKK複合体であり、2つのキナーゼと調節蛋白であるNF-κB essential molecule (NEMO)から構成されるIKKはCARMA1/Bcl-10/MALT-1を介してNEMOのユビキチン化、IKKαまたはIKKβのリン酸化を通じたキナーゼ活性の上昇によりIκBのリン酸化に結びついていく<sup>10)</sup>。シグナルを制御する

分子を各ステップで解除していくことにより、NF-κBの活性化が開始される。逆に言えば、NF-κBはいくつもの制御システムにより精巧に調節されていると言ってよい。ひとたび、NF-κB分子が核内に移行するとT細胞の機能にきわめて重要な遺伝子の転写調節が進行するのである。

### NF-κBの核内移行

NF-κB分子が核内移行すると標的遺伝子のプロモーター領域に存在するκB結合部位にNF-κB分子が直接結合することにより、その遺伝子の転写活性が開始される。5つのNF-κBサブユニットの中で実際に標的遺伝子の転写を司るのはC末に転写活性領域を有するRelA(p65)、RelB、c-Relである(図1)。一方、NF-κB1(p105/p50)とNF-κB2(p100/p52)には転写活性領域は存在しないが、核内移行に重要なシグナルモチーフを有していることから、NF-κB1およびNF-κB2は他の3つのサブユニットとのヘテロダイマーの形成により核内移行を最終的に調節していると考えられる<sup>3)</sup>。NF-κBはT細胞の活性化や増殖に重要なIL-2およびその受容体の一つであるCD25などのようなサイトカインやサイトカイン受容体、ケ

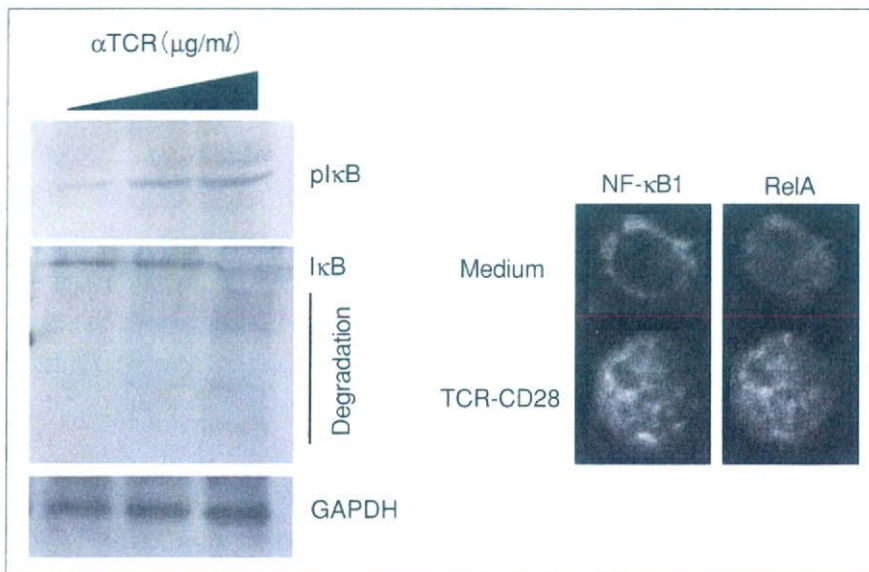


図3 NF-κBの活性化

左：正常マウス脾臓からのCD4陽性細胞に抗TCR抗体(0~1 μg/ml)および抗CD28抗体(20 μg/ml)で刺激し、抗リン酸化IκB抗体によりウエスタンブロットを行った。さらに、IκBの断片化を確認した。ローディングコントロールとしてglyceraldehyde phosphate dehydrogenase (GAPDH)を用いた。  
 右：刺激後のCD4陽性T細胞のNF-κB1およびRelAの核内移行をコンフォーカル顕微鏡解析により確認した。



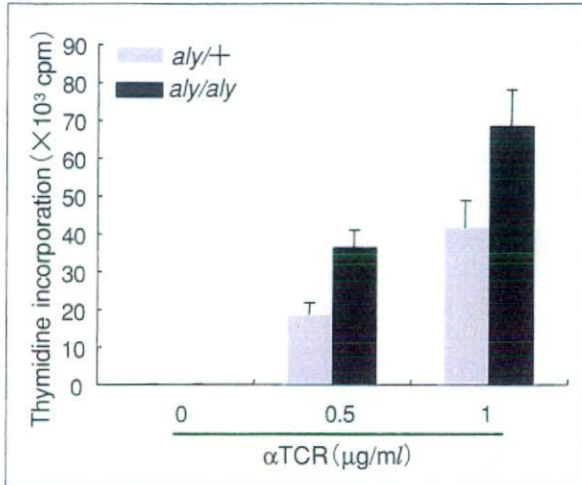


図4 NF-κB2不全マウスのナイーブT細胞の反応性 *aly/aly*マウスと対照マウスの脾臓CD4陽性細胞に抗TCR抗体(0~1 µg/ml)および抗CD28抗体(20 µg/ml)で48時間刺激し、培養の最終12時間における<sup>3</sup>H標識チミジンの取り込みをシンチレーションカウンターにより計測した。

モカイン、接着分子、アポトーシス関連因子、細胞周期関連因子などT細胞の分化、維持および機能獲得に重要な遺伝子群を標的としている<sup>11)</sup>。さらに、NF-κB遺伝子自体をNF-κBが転写調節していることが知られており、転写因子自体の発現調節を自身で制御しているという点はNF-κBそのものの動態を知る上で重要である<sup>12)</sup>。また、各サブユニットがどの標的遺伝子の転写調節に関連しているのかの区別は明確にできていない。各サブユニットの組み合わせや、シグナルカスケードの各中継ポイントにおけるシグナルの質的および量的な違いなど複雑な因子の影響によりNF-κBの活性システムが調節されていることが想定される。

### T細胞における非古典的経路

では、T細胞の活性化に非古典的NF-κB経路は関与してないのであろうか。NF-κB2およびRelBの遺伝子欠損マウスでもT細胞に機能不全や自己免疫病変の発症が報告されている<sup>13)14)</sup>。

正常マウスのCD4陽性T細胞への抗TCR抗体および抗CD28抗体による活性化におけるNF-κBの動態を観察すると、活性の初期段階ではNF-κB1/RelAのヘテロダイマーの核内移行が中心であり、非古典的経路であるNF-κB2/RelBの核内移行はほとんど認められず、増殖期を含む後期段階で

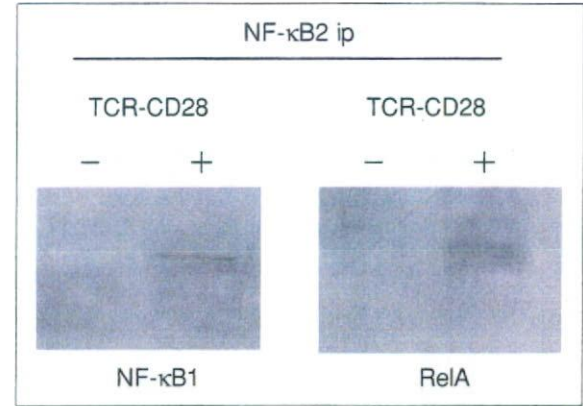


図5 NF-κB2分子とNF-κB1およびRelAとの結合 正常マウス脾臓からのCD4陽性細胞に抗TCR抗体(1 µg/ml)および抗CD28抗体(20 µg/ml)で刺激し、細胞質分画蛋白質を抽出後、NF-κB2抗体で免疫沈降(ip)を行い、ウエスタンブロット法によりNF-κB1およびRelAを検出した。

はNF-κB2/RelBの核内移行が確認されるが<sup>15)</sup>、その役割は不明であった。非古典的NF-κB経路ではTCRおよびCD28からのシグナル伝達に関しては古典的経路ほど明らかにされていない。PKCθ、PI3KおよびAKTからNF-κB-inducing kinase(NIK)を活性化し、NIKはIKKαをリン酸化することによりNF-κB2(p100)のIκB様配列をリン酸化、ユビキチン化する。古典的NF-κB経路と同様にプロテアソームにより断片化され、p52にプロセッシングされることによりp52/RelBヘテロダイマーが核内に移行することが知られている<sup>3)</sup>。NIK遺伝子の点変異マウスである*aly/aly*マウスは末梢CD4陽性T細胞の抗CD3抗体による増殖反応が対照マウスに比較して低下していることが報告されているが<sup>16)</sup>、CD4陽性T細胞の中で、メモリー型のT細胞を除去し、ナイーブ型のCD4陽性T細胞のみを抗TCRおよび抗CD28抗体で刺激して増殖反応を観察すると、対照マウスに比較して有意に高い反応性が認められた(図4)。一方で、*aly/aly*マウスのメモリー型のみ増殖反応はきわめて低くなっていることが判明した<sup>15)</sup>。NIK欠損マウスにおいてもナイーブ型CD4陽性T細胞の増殖活性は有意に上昇していることを確認した。さらに、NF-κB2欠損マウスにおいてもT細胞の過剰増殖が報告されていることから、ナイーブCD4陽性T細胞の活性化における非古典的NF-κB経路(NF-κB2/RelB)の負の制御機構が存



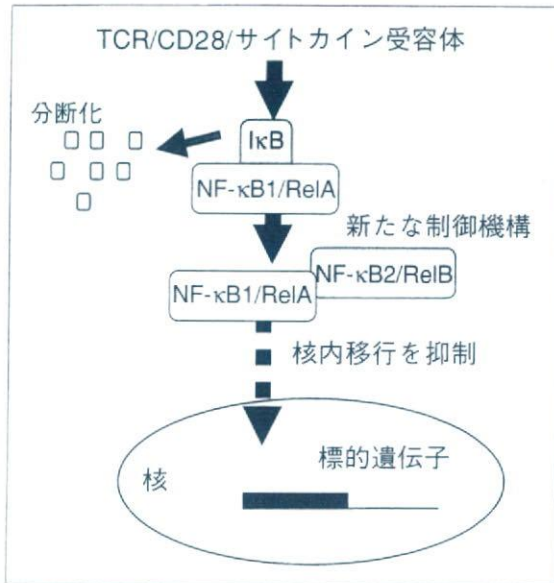


図6 NF-κB2による古典的NF-κB経路の新たな制御機構

在することが明らかになった。また、T細胞のナイーブからメモリー型に至る段階でNF-κBのサブユニットの役割が分担されている可能性が示された。

### NF-κBサブユニット間のクロストーク

NF-κBのサブユニット間での結合に関しては、RelA, RelBおよびc-Relでのホモダイマー以外は5つのサブユニット間でヘテロダイマー、ホモダイマーを形成しうることが知られているが、それらのT細胞における生理的および機能的役割は明らかにされていなかった。上述のNF-κB2の制御機構の細胞内での機序を知るために、正常マウスからナイーブ型(CD44<sup>low</sup>)のCD4陽性T細胞を用いて抗TCRおよび抗CD28抗体で刺激した後、細胞質分画の蛋白質を抽出し、NF-κB2抗体で免疫沈降して各NF-κBサブユニットに対する抗体でウエスタンブロット法により各サブユニットとNF-κB2の直接の結合を確認したところ、NF-κB1およびRelA蛋白が検出された(図5)。一方、*aly/aly*マウスではナイーブ型CD4陽性細胞に刺激を加えると、NF-κB2の合成が阻害されることも判明し、NF-κB2とNF-κB1およびRelAとの結合は観察されなかった<sup>15)</sup>。この所見は*aly/aly*マウスのナイーブ型CD4陽性T細胞の過剰な反応性がNF-κB2分子を介した制御機構の異常により生

じていることを示唆している。従来のIκBによるNF-κBの活性化制御機構に加え、NF-κB2分子がNF-κB1/RelA複合体と細胞質で直接結合することにより、その複合体の核内移行を阻害することでNF-κBの活性化シグナルを調節する新たな免疫制御機構が明らかにされた(図6)。一方で、メモリー型(CD44<sup>high</sup>)CD4陽性T細胞に関してはこの制御システムは働いていないことから、T細胞の活性化段階でNF-κBの各サブユニットの役割が異なっている可能性が示された。非古典的NF-κB経路と古典的NF-κB経路のクロストークはT細胞のみではなく、たとえば、破骨細胞の活性化過程においても報告されていることから、別の細胞種においても、このシステムがなんらかの機能調節に関与している可能性がある<sup>12)17)</sup>。

### NF-κBと免疫疾患

これまでにNF-κB各サブユニットの遺伝子欠損マウスの解析からT細胞におけるNF-κBの生体内での役割が明らかにされている。たとえば、NF-κB1欠損マウスのT細胞では増殖の抑制およびTh<sub>2</sub>タイプのサイトカインの産生減少が観察され、ヒト多発性硬化症の疾患モデル(EAE)での病態感受性の上昇および*Leishmania major*への易感染性がみられ、また、RelA欠損マウスは胎生致死であるが、胎児肝細胞キメラを用いた実験でT細胞の機能低下が報告されている<sup>18)~22)</sup>。一方、RelB欠損マウスは2~3か月齢で重度の自己免疫性貧血および全身の炎症で死亡し、T細胞の機能低下も観察されている<sup>13)</sup>。さらに、NF-κB2欠損マウスではT細胞の過反応がみられ、c-Rel欠損マウスではT細胞の反応性の低下やEAEに対する疾患感受性の低下が知られている<sup>23)24)</sup>。加えて、NF-κB2分子を調節する因子であるNIKの欠損マウスおよびNIK遺伝子の変異マウスにおいてT細胞の増殖は抑制され、多臓器に自己免疫病変が発症することが報告されている<sup>25)</sup>。しかし、T細胞の各活性化段階でのNF-κBの詳細な役割や各サブユニット同士の相互作用と自己免疫との関係は不明な点が多かった。

*aly/aly*マウスやNIK欠損マウスにおける自己免疫病変の発症には胸腺由来のCD25<sup>+</sup>CD4<sup>+</sup>調節性T細胞の数的な減少および質的な障害が自己免



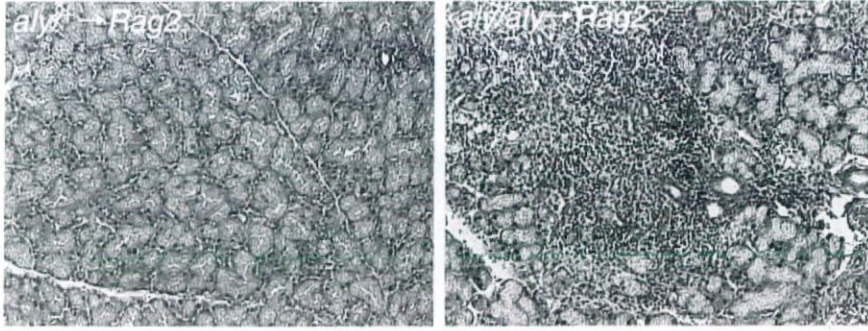


図7 NF- $\kappa$ B2機能不全と自己免疫

*aly/aly*マウスと対照マウスのナイーブ型CD4陽性T細胞を*Rag2*遺伝子欠損マウスに移入後6週における涙腺の病理組織像を示す。*aly/aly*→*Rag2*<sup>-/-</sup>群では腺組織の破壊を伴ったリンパ球浸潤が認められる。

疫の発症に重要であることが報告されている<sup>16)26)</sup>。しかし、*aly/aly*マウスのナイーブ型CD25<sup>-</sup>CD4<sup>+</sup>T細胞を*Rag2*遺伝子欠損マウスに移入すると、本来の*aly/aly*マウスに発症する自己免疫病変よりもはるかに激しい自己免疫性病変が肺や涙腺に認められたことから、調節性T細胞の制御異常に加えて、NF- $\kappa$ B2によるT細胞の制御機構に障害をきたすと自己免疫疾患の発症に結びつくことが示された(図7)。

### まとめ

Nuclear factor (NF)- $\kappa$ Bによる標的遺伝子の転写活性システムはT細胞にとって必須であるといえる。しかし、5つのサブユニットでどれが重要であるのかはいまだ解明されていない。NF- $\kappa$ Bのサブユニット遺伝子のそれぞれの欠損マウスの中にはT細胞の分化や末梢トレランスは正常に維持され、T細胞になんらかの刺激が加えられたときにのみ特定の機能に異常が生じている場合がある。また、なんら機能的異常が見出せない場合も報告されている。サブユニットのどれかが欠損しても別のサブユニットが代替的に機能していることも考えられる。実際の免疫反応においてもT細胞にとってNF- $\kappa$ Bシグナルに障害が生じれば、別のサブユニットが動員され、過剰なシグナルが伝われば各サブユニットを介して制御的に働くNF- $\kappa$ Bネットワークシステムが免疫トレランスの維持に働いているのではなかろうか。今後、NF- $\kappa$ Bシグナルと免疫疾患との関係を明らかにすることは臨床的にもきわめて有用であるといえる。

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## Regulation of Naive T Cell Function By the NF- $\kappa$ B2 Pathway

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Jonathan Sprent

Reprinted from *Nature Immunology*, Volume 7, July 2006





## Regulation of naive T cell function by the NF- $\kappa$ B2 pathway

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T cell activation involves the orchestration of several signaling pathways, including that of the 'classical' transcription factor NF- $\kappa$ B components NF- $\kappa$ B1–RelA. The function of the 'nonclassical' NF- $\kappa$ B2–RelB pathway is less clear, although T cells lacking components of this pathway have activation defects. Here we show that mice deficient in NF- $\kappa$ B-inducing kinase have a complex phenotype consisting of immunosuppression mediated by CD25<sup>+</sup>Foxp3<sup>+</sup> memory CD4<sup>+</sup> cells and, in the absence of those cells, hyper-responsive naive CD4<sup>+</sup> T cells, which caused autoimmune lesions after adoptive transfer into hosts deficient in recombination-activating genes. Biochemical studies indicated involvement of a cell-intrinsic mechanism in which NF- $\kappa$ B2 (p100) limits nuclear translocation of NF- $\kappa$ B1–RelA and thereby functions as a regulatory 'brake' for the activation of naive T cells.

The transcription factor NF- $\kappa$ B is key in the regulation of many inflammatory processes of immune cells<sup>1</sup>. The NF- $\kappa$ B family consists of five subunits: NF- $\kappa$ B1 (p105-p50), NF- $\kappa$ B2 (p100-p52), RelA (p65), RelB and c-Rel. Hetero- or homodimers of these subunits can be translocated into the nucleus to bind to  $\kappa$ B sequences of neighboring 'target' genes, thus regulating the transcription of genes required for cell activation, survival and development<sup>2</sup>. Two pathways of NF- $\kappa$ B have been defined in immune cells<sup>3</sup>: the 'classical' pathway, which is initiated by complexes of NF- $\kappa$ B1 and RelA, and an alternative or 'nonclassical' pathway, which is initiated by complexes of NF- $\kappa$ B2 and RelB.

For T cells, stimulation via the T cell receptor (TCR) and costimulatory molecules such as CD28 leads to NF- $\kappa$ B activation through a variety of intracellular signaling molecules<sup>4</sup>. Initially, TCR-CD28 signaling via many adaptor molecules leads to activation of protein kinase C- $\theta$  (PKC- $\theta$ )<sup>5</sup>. Thereafter, CARMA1–Bcl-10–MALT1 proteins 'downstream' of PKC- $\theta$  activate I $\kappa$ B kinase (IKK) complexes, including IKK $\alpha$ , IKK $\beta$  and the adaptor protein IKK $\gamma$  (also called NEMO)<sup>6,7</sup>. Activated IKK complexes then phosphorylate I $\kappa$ B, releasing it from its constitutively bound state with cytoplasmic NF- $\kappa$ B complexes (mainly NF- $\kappa$ B1–RelA) that normally prevents NF- $\kappa$ B complexes from translocating to the nucleus. Ubiquitination and degradation of I $\kappa$ B by IKK complexes allows components of the classical NF- $\kappa$ B pathway, especially p50–RelA, to be transported into the nucleus, thus promoting transcription of essential target genes required for survival, cytokine and chemokine production, upregulation of adhesion molecules, organogenesis and apoptosis in the immune system<sup>8</sup>. The classical NF- $\kappa$ B pathway is especially important for the synthesis of interleukin 2 (IL-2) as well as IL-2 receptor (IL-2R, also called CD25) in T cells<sup>9,10</sup>.

As for the alternative, nonclassical NF- $\kappa$ B pathway, signals from specific cytokine receptors (such as the lymphotoxin- $\beta$  receptor) activate NF- $\kappa$ B-inducing kinase (NIK) as well as PKC- $\theta$  and CARMA1–Bcl-10–MALT1, which in turn activate homodimers of IKK $\alpha$  that are required for the cleavage of p100 to p52 (refs. 11–13). Complexes of p52 and RelB then translocate into the nucleus to regulate transcription of a different set of genes.

Studies of gene-knockout mice have demonstrated that individual members of the NF- $\kappa$ B family have distinct roles *in vivo* in T cell function. Thus, T cell proliferation and T helper type 2 cytokine production is reduced in NF- $\kappa$ B1-deficient (*Nfkb1*<sup>-/-</sup>) mice, and these mice show increased susceptibility to experimental autoimmune encephalomyelitis, typhlocolitis and infection with *Leishmania major* but are resistant to asthma<sup>14–17</sup>. RelA-deficient (*Rela*<sup>-/-</sup>) mice have a phenotype that is embryonically lethal, but studies with fetal liver chimeras indicate that *Rela*<sup>-/-</sup> T cells are functionally defective<sup>18,19</sup>. RelB-deficient (*Relb*<sup>-/-</sup>) mice are viable but develop systemic inflammation and severe anemia around 2–3 months of age, and the T cell and B cell functions of these mice are suppressed<sup>20</sup>. NF- $\kappa$ B2-deficient (*Nfkb2*<sup>-/-</sup>) mice have B cell defects as well as T cell hyperplasia and hyperactivation of dendritic cells<sup>21,22</sup>. Finally, mice deficient in c-Rel (*Rel*<sup>-/-</sup>) have B cell defects, impaired T cell proliferative responses and reduced susceptibility to experimental autoimmune encephalomyelitis<sup>23,24</sup>. Nevertheless, precise information about the functions of the individual NF- $\kappa$ B family members on T cell function is still unclear.

The importance of NIK for NF- $\kappa$ B activation has been demonstrated in studies of NIK-deficient (*Map3k14*<sup>-/-</sup>) mice and also mice with alymphoplasia (*Map3k14*<sup>aly/aly</sup> or 'aly/aly' mice), which carry a mutation of *Map3k14*. Functionally, NIK is key in regulating the

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Received 27 January; accepted 26 April; published online 28 May 2006; doi:10.1038/ni1351