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Complementary Signaling through flt3 and Interleukin-7 Receptor α Is Indispensable for Fetal and Adult B Cell Genesis

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Abstract

Extensive studies of mice deficient in one or several cytokine receptors have failed to support an indispensable role of cytokines in development of multiple blood cell lineages. Whereas B1 B cells and Igs are sustained at normal levels throughout life of mice deficient in IL-7, IL-7R α , common cytokine receptor gamma chain, or flt3 ligand (FL), we report here that adult mice double deficient in IL-7R α and FL completely lack visible LNs, conventional IgM⁺ B cells, IgA⁺ plasma cells, and B1 cells, and consequently produce no Igs. All stages of committed B cell progenitors are undetectable in FL^{-/-} $\times$ IL-7R α ^{-/-} BM that also lacks expression of the B cell commitment factor *Pax5* and its direct target genes. Furthermore, in contrast to IL-7R α ^{-/-} mice, FL^{-/-} $\times$ IL-7R α ^{-/-} mice also lack mature B cells and detectable committed B cell progenitors during fetal development. Thus, signaling through the cytokine tyrosine kinase receptor flt3 and IL-7R α are indispensable for fetal and adult B cell development.

Key words: lymphopoiesis • IL-7 receptor • Flt3 ligand • Pax5 • B1 cells

Introduction

Hematopoiesis is a tightly regulated process in which multipotent hematopoietic stem cells commit and differentiate along a number of specific pathways to continuously replenish cells of all blood lineages (1). Several lineage-restricted hematopoietic growth factors (cytokines) and their corresponding receptors promote the development of specific blood cell lineages in vitro and in vivo (2–5). However, targeted deletions of most of these cytokine receptors or their ligands have resulted in surprisingly mild phenotypes and rarely in complete losses of specific blood cell lineages (3, 6–12). Furthermore, since the physiological functions of

most blood lineages remain intact at reduced cell numbers, such mutations frequently have limited or no functional consequences. For instance, although thrombopoietin-deficient mice have an 85% reduction in platelets, they have no bleeding disorder (13).

As a result of these observations, the concept and controversy of cytokine redundancy has emerged, implicating that many lineage-restricted cytokines may play a limited role in the regulation of hematopoiesis. This latter point also raises the possibility that combinatorial interactions between two or more cytokines may frequently be required for development of specific blood cell lineages (3). The last possibility

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Abbreviations used in this paper: AP, alkaline phosphatase; CLP, common lymphoid progenitor; FL, flt3 ligand; PB, peripheral blood; γ c, γ receptor chain.

has already provoked extensive research but so far provided limited evidence for the requirement of such in vivo synergy between lineage-restricted cytokines and, for instance, in myeloid development failed to support an absolute combinatorial cytokine requirement for the formation of both platelets and granulocytes (14–24).

Several cytokines have been implicated to be involved in the regulation of B cell development (25). Of these, IL-7 has emerged as a key cytokine involved in B cell genesis (9, 26, 27). However, recent studies have demonstrated that although B cell development is almost completely arrested at a very early stage in the BM of adult IL-7 and IL-7R α -deficient mice, a pool of mature B cells, primarily of fetal and perinatal origin, is sustained and stable throughout adult life, capable of producing normal levels of Igs (28–30). Furthermore, the compartment of self-replenishing B1 cells is normal in IL-7-deficient mice and has therefore been suggested to be IL-7 independent (28).

Whether IL-7 in combination with other cytokines might be indispensable for B cell genesis has been thoroughly investigated in mice with targeted deletions of the common γ receptor chain (γ c), essential for signaling by six lymphokines (IL-2, IL-4, IL-7, IL-9, IL-15, and IL-21). However, γ c^{-/-} mice have a B cell phenotype corresponding closely to that of IL-7R α ^{-/-} mice (31, 32), suggesting that among known lymphokines, only IL-7 is critically involved in B cell genesis and that the other lymphokines using γ c may not be required for B cell development.

Unlike lineage-restricted cytokine receptors belonging to the hematopoietin receptor superfamily (33), the fms-like tyrosine kinase-3 (flt3 or flk2) receptor is expressed primarily at early stages of hematopoiesis and in particular early lymphoid development (34). Although flt3 expression is lost at the pre-B cell stage (35) and flt3 ligand (FL)-deficient mice only have marginally reduced mature B cell numbers and normal Ig levels (36, 37), there is ample evidence that flt3 and its ligand indeed play an important role in B cell development. Most importantly, the BM of flk2 and FL-deficient mice have reduced levels of B cell progenitors (36, 38), including the common lymphoid progenitor (CLP) (37). Furthermore, the combined action of FL and IL-7 is both sufficient and strictly required for stroma-independent in vitro B cell development from adult multipotent hematopoietic progenitors (39).

Thus, we hypothesized that rather than a combination of lymphokines, the concerted action of two cytokine receptors—flt3, an early acting tyrosine kinase receptor and the later acting hematopoietin receptor IL-7R α —might be indispensable for B cell genesis in vivo. Herein we present compelling evidence supporting this hypothesis. During both fetal and adult hematopoiesis, mice deficient in the expression of FL and IL-7R α completely lack mature conventional IgM⁺ B cells, IgA⁺ plasma cells, and B1 B cells, and in striking contrast to FL or IL-7R α single-deficient mice, fail to produce circulating Ig.

Materials and Methods

Generation of Double Knockout Mice Deficient in FL and IL-7 α Expression. Mice deficient in FL (36) or IL-7R α (26) expression were generated as described previously. FL^{-/-} mice were on a pure C57BL/6 background (36), whereas IL-7R α ^{-/-} mice had been backcrossed for five generations with C57BL/6 mice. Mice double deficient for FL and IL-7R α expression were obtained by crossbreeding of single FL^{-/-} and IL-7R α ^{-/-} mice and subsequent interbreeding of heterozygous FL^{+/-} IL-7R α ^{+/-} mice. Mice used for experiments were obtained through breeding of FL^{-/-} $\times$ IL-7R α ^{-/-} mice, and C57BL/6 mice were used as WT controls. All mice used for the study were genotyped using PCR with specific primers to verify the absence of FL and IL-7R α genes (26, 36). All studies of adult B lymphopoiesis were performed on age-matched mice, 8–12 wk old, kept under specific pathogen-free conditions.

For analysis of embryonic development, timed pregnancies were determined as described previously (40); the day the vaginal plug was observed was considered as day 0. All performed experiments were approved by the Ethical Committee at Lund University.

Tissues. Peripheral blood (PB) was collected from the retro-orbital sinus venous plexus or vena cava into heparinized tubes. White blood cells were isolated using 1% dextran (Amersham Biosciences) solution, and the remaining red cells were lysed using ammonium chloride. BM cell suspensions were prepared by flushing isolated tibiae and femora with PBS (Sigma-Aldrich) containing 5% FCS (BioWhittaker). Cell suspensions were prepared from spleens, mesenteric LNs, and Peyer's patches by gently breaking up the tissues in PBS (Sigma-Aldrich) containing 5% FCS. Lamina propria lymphocytes were isolated as described before (41). Spleens and livers were collected from embryos at day 17–18 of gestation, and cell suspensions were prepared by gently breaking up the tissues in PBS (Sigma-Aldrich) containing 5% FCS.

mAbs and Flow Cytometry Analysis. The following antibodies were used for flow cytometry: anti-B220 (RA3-6B2), anti-CD19 (1D3), anti-CD5 (53-7.3), anti-CD43 (S7), anti-NK1.1 (PK136), anti-IgM (R6-60.2), anti-CD3e (145-2C11), anti-CD11b (M1/70) (all from Amersham Biosciences), and anti-IgA (11-44-2) (Southern Biotechnology Associates, Inc.). Anti-AA4.1 antibody was a gift from Dr. Ihor Lemischka (Princeton University, Princeton, NJ). Isotype-matched controls used included mouse IgG1, mouse IgG2a, rat IgG2a, rat IgG2b, and hamster IgG labeled with appropriate fluorochromes (BD Biosciences). Up to 10⁶ cells per sample were first incubated in PBS (Sigma-Aldrich) containing 5% FCS (BioWhittaker) and Fc block (CD16/CD32; BD Biosciences) to minimize nonspecific binding. Cells were then labeled with the appropriate antibodies for 15 min at 4°C in PBS containing 5% FCS. Finally, 7AAD was added to exclude dead cells from the analysis. Samples were analyzed on FACSCalibur (Becton Dickinson). Between 50,000 and 400,000 events were collected, and analysis was performed using CellQuest (Becton Dickinson) and FlowJo (Tree Star Inc.) software. All FACS® data were displayed using a log 10 scale.

Ig ELISA. ELISA analysis was performed according to Engvall and Perlman (42). Briefly, microtiter ELISA plates (Costar Corning 3590) were plated with 100 μ l/well of 10 μ g/ml catcher antibody in PBS and incubated overnight at 4°C. For blocking of nonspecific binding, the solutions were replaced with 200 μ l/well 1% BSA in PBS and incubated overnight at 4°C. The plates were washed three times with 200 μ l/well washing buffer (0.05% Tween 20, 0.02% NaN3 in PBS) and then incubated overnight at 4°C with different dilutions of the serum

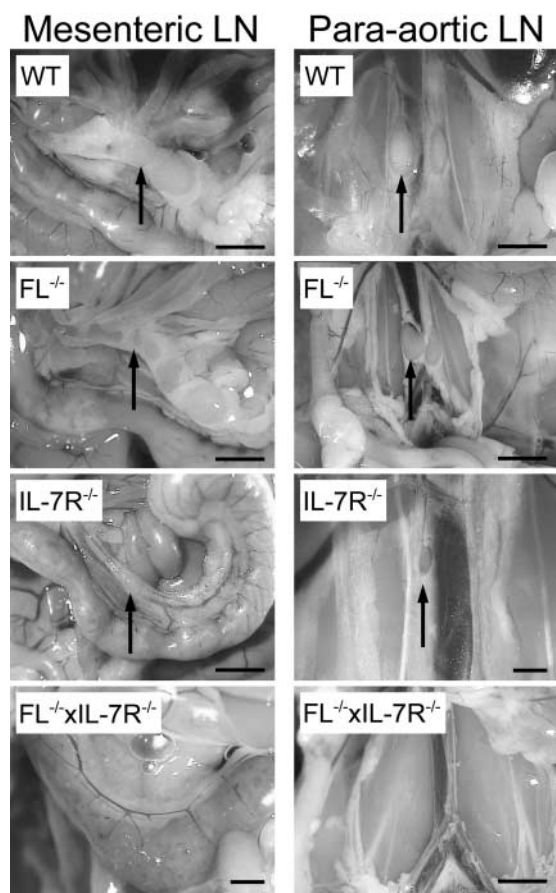


Figure 1. Mice double deficient in *fli3* ligand and IL-7R α expression show impairment in the development of lymphoid tissue. Photographs of mesenteric and para-aortic LNs taken with 2.2 to 3.45 magnification; arrows point to location of LNs. Bar, 2.2 cm.

samples (IgM, IgA: 1:50,00 and 1:20,000; IgG: 1:20,000 and 1:80,000). Serum from FL^{-/-} $\times$ IL-7R α ^{-/-} mice was in addition diluted at 1:100 and 1:400 for all analysis. The plates were washed three times with 200 μ l/well washing buffer and further incu-

bated with 100 μ l/well alkaline phosphatase (AP) coupled detecting antibody (IgM: 1.2 μ g/ml; IgG: 1.2 μ g/ml), or biotinylated anti-IgA (2.5 μ g/ml), overnight at 4°C. Only for detection of IgA, the plates were washed three times with 200 μ l/well washing buffer and incubated overnight at 4°C with 100 μ l/well AP conjugated with streptavidin (1 μ g/ml). After washing three times with 200 μ l/well washing buffer, 100 μ l/well AP developing solution was added (1 M diethanolamine, 4 mM MgCl₂, pH 9.8 with freshly added 0.5 mg/ml pNPP [no. O-104; Sigma-Aldrich]). The reaction was developed at room temperature and measured at several time points on an ELISA reader at 415 nm. The following antibodies were used: donkey anti-mouse IgM (low cross reactivity to rat) (Jackson ImmunoResearch Laboratories), donkey anti-mouse IgM AP (low cross reactivity to rat) (Jackson ImmunoResearch Laboratories), donkey anti-mouse IgG (low cross reactivity to rat) (Jackson ImmunoResearch Laboratories), donkey anti-mouse IgG AP (low cross reactivity to rat) (Jackson ImmunoResearch Laboratories), rat anti-mouse IgA (C10-3; BD Biosciences), biotinylated rat anti-mouse IgA (C10-1; BD Biosciences), and AP-conjugated streptavidin (Jackson ImmunoResearch Laboratories).

D-J Rearrangement Analysis. BM cells were suspended into single cells by gentle mechanic stress and filtering to remove debris. Genomic DNA was prepared using Trizol (GIBCO BRL) according to the manufacturers instructions. D-J rearrangements were then assayed by PCR using a protocol based on that by Schlissel et al. (43). The D-J-rearranged and germline-configured DNAs were amplified in the same reaction by 30 cycles (94°C for 30 s, 60°C for 45 s, and 72°C for 1 min) using the DH and J3 primer at 1 μ M and the Mu0 primer at 0.1 μ M final concentration. The Mu0 amplifies together with the J3 primer germline DNA and the aim of using reduced amounts of Mu0 was to reduce the signal from the germline amplification. The PCR products were blotted onto Hybond N⁺ nylon membranes (Amersham Biosciences) using capillary blotting. Membranes were prehybridized in 5 $\times$ Denhardt's, 6 $\times$ SSPE, 0.1% SDS, and 100 μ g/ml Salmon Sperm DNA at 45°C for 60 min and hybridized with a γ -³²P-labeled oligonucleotide complementary to the J3 region (JH3) for 12 h at 45°C in the same solution. Membranes were washed at room temperature in 2 $\times$ SSC supplemented with 0.1% SDS for 15 min and 0.1 $\times$ SSC with 0.1% SDS for 5–10 min. The hybridized membrane was then subject to autoradiography.

Table I. Mice Double Deficient in *fli3* Ligand and IL-7R α Expression Have Impaired Development of Lymphoid Tissues

Tissue	WT Cellularity $\times 10^6$	FL ^{-/-} % WT control	IL-7R α ^{-/-} % WT control	FL ^{-/-} $\times$ IL-7R α ^{-/-} % WT control
BM	80 (11)	77 (16)	99 (7)	102 (8)
Spleen	187 (39)	60 (14)	22 (6)	16 (3)
PB	6.8 (1.7) (per ml)	73 (24)	25 (3)	10 (3) ^a
^c Mesenteric LNs	17 (3)	78 (19)	6.3 (1.2)	Lacking ^b
^d Peyer's Patches number	7 (1)	4 (2)	2 (1)	Lacking ^b
Peyer's Patches total cellularity	1.7 (0.6)	53 (16)	5.5 (1.5)	

All analysis of double knockout mice lacking FL and IL-7R α expression (FL^{-/-} $\times$ IL-7R α ^{-/-}) was done in parallel with age-matched 8–12-wk-old WT control and single FL^{-/-} and IL-7R α ^{-/-} mice. Data are expressed as means (SD) from 6–12 mice analyzed in each group.

^{a,b}Statistically significant differences between single IL-7R α ^{-/-} and FL^{-/-} $\times$ IL-7R α ^{-/-} mice, with P < 0.01 and P < 0.05, respectively.

^cTotal cellularity in mesenteric LNs.

^dNumber of Peyer's Patches; mean (SD) values from WT, FL^{-/-}, IL-7R α ^{-/-}, and FL^{-/-} $\times$ IL-7R α ^{-/-} mice.

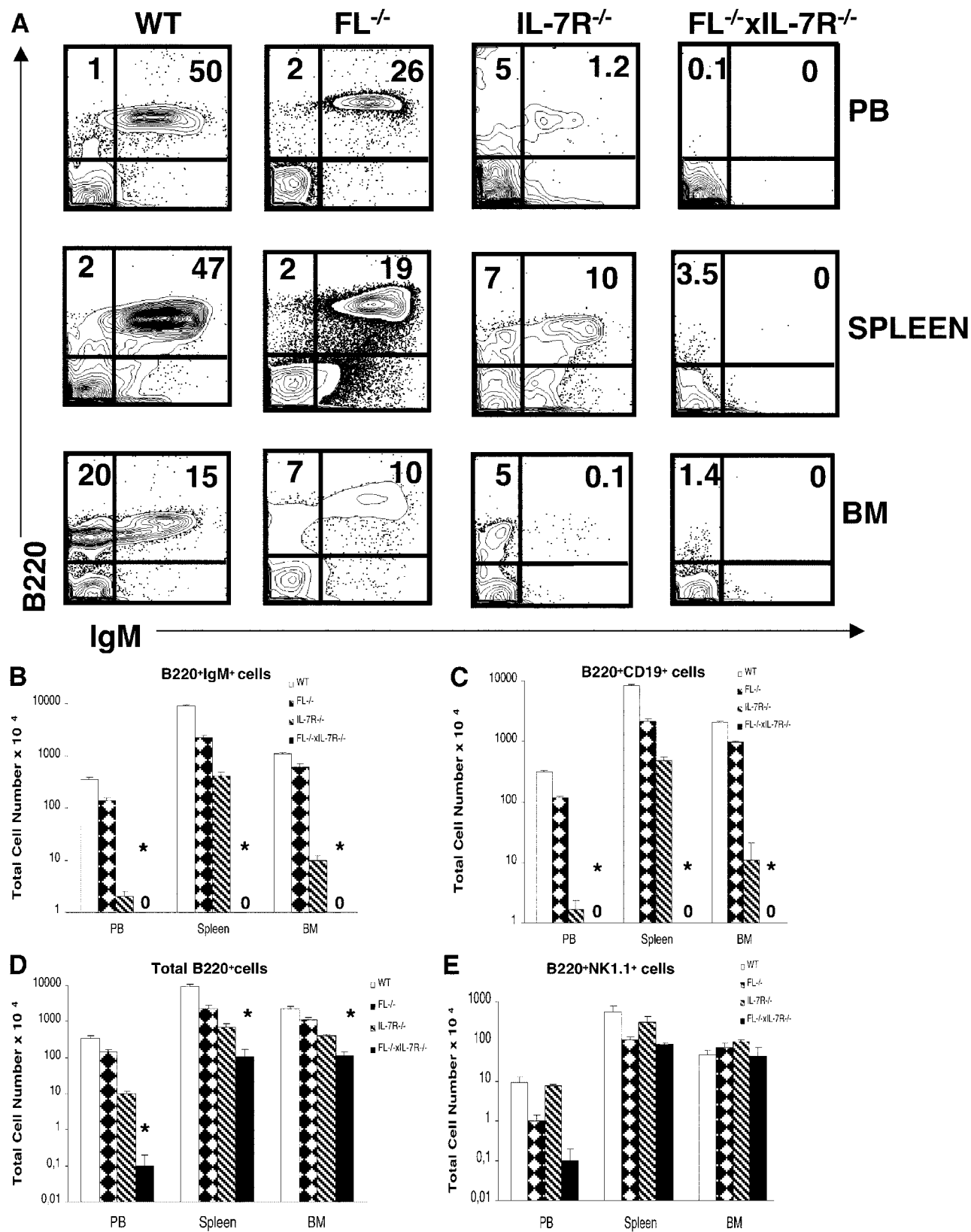


Figure 2. Adult mice double deficient in Flt3 ligand and IL-7R α expression lack B cells in the PB, spleen, and BM. PB, spleen, and BM cells from 8–12-wk-old mice were stained with mAbs against B220, IgM, CD19, and NK1.1 (as described in Materials and Methods). (A) Plots of B220 and IgM expression in WT, FL-deficient, IL-7R α -deficient and FL^{-/-} × IL-7R α ^{-/-} double deficient mice, respectively. Numbers represent mean values from 6–12 mice. (B) Total number of B220⁺ cells coexpressing IgM, B220⁺ cells expressing CD19 (C), B220⁺ cells (D), and B220⁺ cells expressing NK1.1

The following oligonucleotides were used for PCR: DH, 5'-GGAATTCG(A/C)TTTTTGT(C/G)AAGGGATCTACTAC-TGTG-3'; J3, 5'-GTCTAGATTCTCACAAGAGTCCGATA-GACCTTGG-3'; Mu0, 5'-CCGCATGCCAAGGCTAGCCT-GAAAGATTACC-3'. The following oligonucleotides was used for hybridization: JH3, 5'-AGACAGTGACCAGAGTCCCTTGG-3'.

Expression of B Cell Genes. Total RNA was prepared using RNazol (Tel-Test) from 10 million BM cells and dissolved in 50 μ l. 10 μ l was used for cDNA preparation in a total volume of 20 μ l using Superscript II and random primers (Life Technologies). 1 μ l cDNA was used per PCR reaction, except serial fivefold dilutions for HGPRT starting with 1 μ l. The following conditions were used for PCR reactions: HGPRT, 30 cycles at 55°C annealing temperature; all other reactions, 35 cycles at 55–58°C annealing temperature.

The following primer sequences were used: pax-5, sense, 5'-TCCTCGGACCATCAGGACAG-3' and antisense, 5'-CCT-GTTGATGGAGCTGACGC-3'; for mb-1 sense, 5'-GGGGT-CTAGAAGCCCTCAGAGCCCTGCCTC-3' and antisense, 5'-TGCCGAATTCCTGGTAGGTGCCCTGGAGTC-3'; and CD19, sense, 5'-TACCTTAGTAGGAGGCAGGCC-3' and antisense, 5'-CCGGAACATCTCCCCACTAT-3'.

Statistics. All results were expressed as means ($\pm$ SD). The statistical significances between groups were determined using the Student's *t* test.

Results

Mice Double Deficient in *flt3* Ligand and IL-7R α Expression Are Severely Immune Compromised and Lack Peyer's Patches and Identifiable LNs. Initially, FL^{-/-} $\times$ IL-7R α ^{-/-} mice showed reduced survival compared with either FL^{-/-} or IL-7R α ^{-/-} mice. The reduced survival was most likely due to enhanced susceptibility to opportunistic infections, since breeding and housing of FL^{-/-} $\times$ IL-7R α ^{-/-} mice in specific pathogen-free conditions resulted in mice with a normal life span.

No significant differences in BM cellularity were observed between WT, IL-7R α ^{-/-}, FL^{-/-}, or FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Table I). In contrast, spleen cellularity in FL^{-/-}, IL-7R α ^{-/-}, and FL^{-/-} $\times$ IL-7R α ^{-/-} mice was reduced to 60, 22, and 16%, respectively, of WT controls. The total peripheral white blood cell counts in FL^{-/-} $\times$ IL-7R α ^{-/-} mice were more significantly reduced (to 10% of WT mice) compared with either single FL^{-/-} or IL-7R α ^{-/-} mice (Table I). Previous studies have reported reduced cellularity in the LNs and Peyer's Patches of IL-7R α ^{-/-} mice (26, 44). In contrast to IL-7R α ^{-/-} mice in which LNs were present but with reduced cellularity, no peripheral (e.g., para-aortic [Fig. 1], axial, and inguinal [not depicted]) or mesenteric LNs were identifiable in FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Table I and Fig. 1). Similarly, unlike IL-7R α ^{-/-} mice where Peyer's Patches were present at reduced numbers (and cellularity), these could not be detected in FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Table I). Collectively, the se-

vere lymphoid hypoplasia detected in FL^{-/-} $\times$ IL-7R α ^{-/-} mice suggests a critical and complimentary role of *flt3* and IL-7R α signaling in lymphoid development.

Absence of Mature IgM⁺ B Cells in the Blood, Hematopoietic, and Lymphoid Tissues of Adult FL^{-/-} $\times$ IL-7R α ^{-/-} Mice. The absence of identifiable LNs in FL^{-/-} $\times$ IL-7R α ^{-/-} mice suggested that the generation of mature B cells may be more severely affected in FL^{-/-} $\times$ IL-7R α ^{-/-} than either FL^{-/-} or IL-7R α ^{-/-} mice. Thus, the presence of mature B220⁺IgM⁺ B cells in the PB, spleen, and BM of 8–12-wk-old FL^{-/-} $\times$ IL-7R α ^{-/-} mice was next investigated (Fig. 2, A and B). As reported previously (36), B220⁺IgM⁺ cells were reduced by 53–75% in FL^{-/-} mice (Fig. 2, A and B), whereas IL-7R α ^{-/-} mice had clearly reduced but detectable B220⁺IgM⁺ cells in PB, BM, and spleen in agreement with previous studies (26) (Fig. 2, A and B). In striking contrast, FL^{-/-} $\times$ IL-7R α ^{-/-} mice completely lacked mature B220⁺IgM⁺ B cells in the PB, spleen, and BM (Fig. 2, A and B).

The absence of B cells in FL^{-/-} $\times$ IL-7R α ^{-/-} mice was further confirmed by a complete lack of B220⁺CD19⁺ cells in PB, spleens, and BM, whereas these were present in IL-7R α ^{-/-} mice (Fig. 2 C). Although present at dramatically reduced levels compared with FL^{-/-} and IL-7R α ^{-/-} mice, B220⁺ cells were detectable in the PB, spleen, and BM of FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Fig. 2 D). Noteworthy, however, most of the rare B220⁺ cells detected in FL^{-/-} $\times$ IL-7R α ^{-/-} mice coexpressed NK1.1 (Fig. 2 E), consistent with the phenotype of NK cell progenitors rather than bona fide B cells (45). Thus, in contrast to either FL^{-/-} or IL-7R α ^{-/-} mice, adult FL^{-/-} $\times$ IL-7R α ^{-/-} mice completely lack B cells.

In contrast to the complete absence of mature B cells, the number of PB T (CD3⁺) cells was only slightly reduced in FL^{-/-} $\times$ IL-7R α ^{-/-} mice compared with IL-7R α ^{-/-} mice (Table II), whereas the number of myeloid (CD11b⁺) cells was comparable to that of FL^{-/-} mice (Table II).

Absence of IgA⁺ Plasma Cells in Lamina Propria and Peritoneal B1 Cells in Mice Double Deficient in *flt3* Ligand and IL-7R α Expression. Since production of IgA has been demonstrated to be regulated by mechanisms distinct from those of IgM (46), FL^{-/-} $\times$ IL-7R α ^{-/-} mice were next examined for the presence of IgA⁺ plasma cells in the lamina propria of the gut (Fig. 3 A).

The number of lymphocytes in the lamina propria was reduced by 79 $\pm$ 4% in FL^{-/-} $\times$ IL-7R α ^{-/-} mice, whereas only a slight reduction in lymphocyte numbers was detected in the lamina propria of FL^{-/-} and IL-7R α ^{-/-} mice (14 $\pm$ 12% and 34 $\pm$ 4% reduction, respectively). Strikingly, whereas IgA⁺ plasma cells were present in normal numbers in FL^{-/-} mice and reduced by 75% in IL-7R α ^{-/-} mice, IgA⁺ plasma cells were completely absent in FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Fig. 3 A).

(E). BM cellularity represents cell counts from both tibiae and femora, whereas total number of white blood cells in PB was calculated per 1 ml. Data in B–E are expressed as mean values (SD) from 6–12 age-matched mice in each group. *Statistically significant differences (*P* < 0.01) between IL-7R α ^{-/-} and FL^{-/-} $\times$ IL-7R α ^{-/-} mice. 0, no B220⁺IgM⁺ or B220⁺CD19⁺ cells were detected in FL^{-/-} $\times$ IL-7R α ^{-/-} mice.

Table II. Presence of T and Myeloid Cells in Mice Double Deficient in *flt3* Ligand and IL-7R α Expression

Mouse	Total number of T (CD3 ⁺) cells per 1 ml $\times 10^4$	Total number of myeloid (CD11b ⁺) cells per 1 ml $\times 10^4$
WT	163 (47)	100 (30)
FL ^{-/-}	220 (13)	73 (21)
IL-7R α ^{-/-}	21 (8)	120 (29)
FL ^{-/-} $\times$ IL-7R α ^{-/-}	12 (6)	70 (16)

Cells from peripheral blood from 8–12-wk-old mice were stained with mAbs against CD3 and CD11b (described in Materials and Methods). Data represent total number of cells and are expressed as mean values (SD) from four age-matched mice in each group.

B1a cells, characterized by expression of low levels of B220 and intermediate levels of CD5, represent a subset of self-replenishing B cells that arise early during fetal development before the appearance of conventional B cells (47,

48). Several studies suggest that B1 cells are regulated by mechanisms distinct from those of conventional B cells. This is supported by studies of $\gamma c^{-/-}$ and IL-7^{-/-} mice in which conventional B cells are dramatically reduced, whereas B1 cells are present at near normal levels (28, 31). Whereas the number of peritoneal leukocytes was normal ($106 \pm 4\%$) in FL^{-/-} mice and present at almost normal levels ($70 \pm 10\%$) in IL-7R α ^{-/-} mice, they were significantly reduced in FL^{-/-} $\times$ IL-7R α ^{-/-} mice ($27 \pm 4\%$; $P < 0.01$ compared with IL-7R α ^{-/-}). Whereas the total number of B1a cells was normal in FL^{-/-} mice and reduced by 63% compared to control levels in IL-7R α ^{-/-} mice, they were completely absent in FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Fig. 3 B). Thus, signaling through *flt3* and IL-7R α is also critical for generation and/or maintenance of B1a cells.

*Mice Double Deficient in *flt3* ligand and IL-7R α Expression Fail To Produce Igs.* Although $\gamma c^{-/-}$ mice and IL-7R α ^{-/-} mice have reduced numbers of conventional B cells, these mice have normal levels of circulating Ig (30, 31). This could reflect that residual numbers of conventional B cells and/or normal numbers of B1 cells (known to produce all Ig types) (47) are capable of sustaining Ig production in

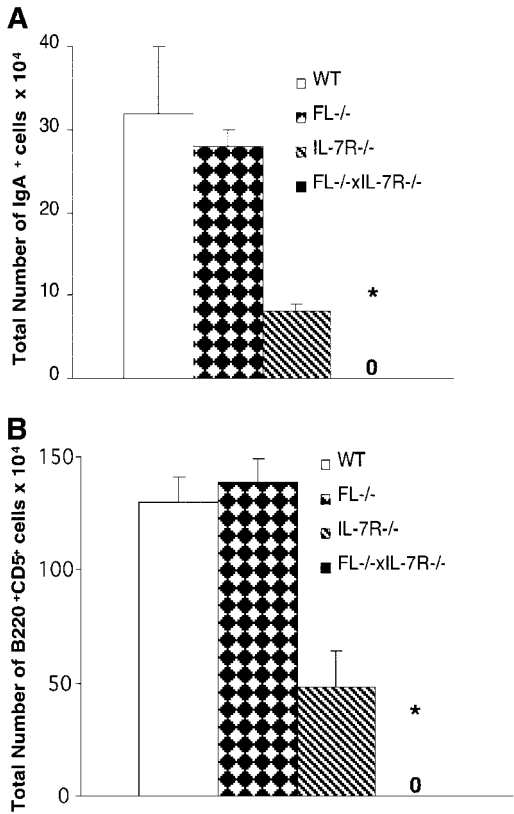


Figure 3. Mice double deficient in *Flt3* ligand and IL-7R α expression have reduced numbers of IgA⁺ plasma cells and B1 cells. (A). Cells from lamina propria of 8–12-wk-old WT, FL^{-/-}, IL-7R α ^{-/-}, and FL^{-/-} $\times$ IL-7R α ^{-/-} mice were stained with mAbs against IgA (as described in Materials and Methods). 0, no IgA⁺ cells were detected in FL^{-/-} $\times$ IL-7R α ^{-/-} mice. (B) Cells from peritoneal lavage of 8–12-wk-old WT, FL^{-/-}, IL-7R α ^{-/-}, and FL^{-/-} $\times$ IL-7R α ^{-/-} mice were stained with mAbs against B220 and CD5 (described in Materials and Methods). All data are expressed as mean values (SD) from three to six mice. *Indicates statistically significant differences ($P < 0.01$) between IL-7R α ^{-/-} and FL^{-/-} $\times$ IL-7R α ^{-/-} mice. 0, no B220⁺CD5⁺ cells were detected in FL^{-/-} $\times$ IL-7R α ^{-/-} mice.

Table III. Mice Double Deficient in *flt3* Ligand and IL-7R α Expression Do Not Produce Igs

Mouse	IgM levels $\mu\text{g/ml}$	IgG levels $\mu\text{g/ml}$	IgA levels $\mu\text{g/ml}$
WT	198 $\pm$ 95	625 $\pm$ 117	186 $\pm$ 84
FL ^{-/-}	Normal ^a	Normal ^a	Normal ^a
IL-7R α ^{-/-}	237 $\pm$ 53	501 $\pm$ 314	39 $\pm$ 21
FL ^{-/-} $\times$ IL-7R α ^{-/-}	0 ^b (bd)	0 ^b (bd)	0 ^b (bd)

The serum levels of IgM, IgG, and IgA were analyzed using ELISA (described in Materials and Methods) with the following detection limits: 0.5 $\mu\text{g/ml}$ for IgM, 0.15 $\mu\text{g/ml}$ for IgG, and 0.08 $\mu\text{g/ml}$ for IgA. 6–10 age-matched mice were analyzed per each group, homozygous Rag^{-/-} mice were used as negative controls (66). bd, below detection. ^aReference 36.

^bStatistically significant difference ($P < 0.01$) between IL-7R α ^{-/-} and FL^{-/-} $\times$ IL-7R α ^{-/-} mice.

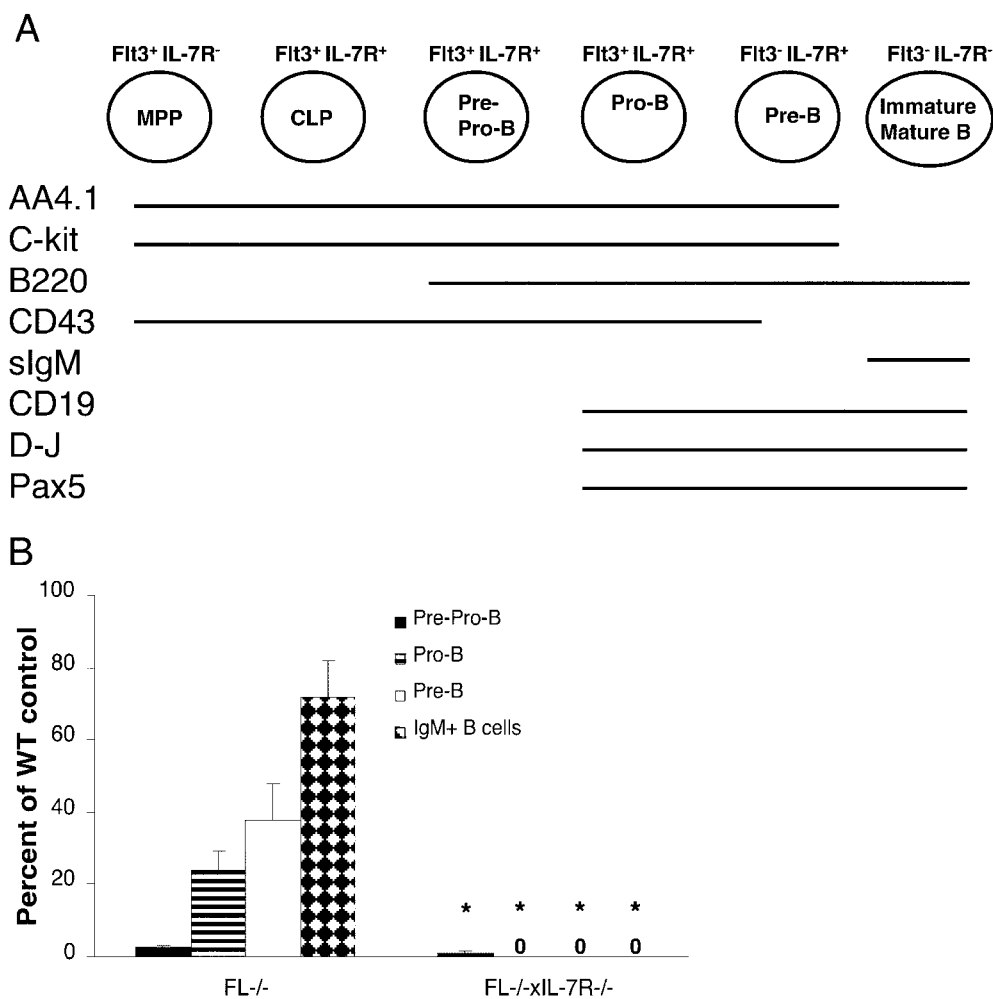


Figure 4. Mice double deficient in flt3 ligand and IL-7R α expression show dramatic reductions in early and late B cell progenitors in adult BM. (A) Schematic illustration of cell surface phenotype, B cell gene expression, and D-J rearrangement during normal B cell development. Also shown are expression patterns for flt3 and IL-7R. MPP, multipotent progenitor, Lin⁻Sca-1⁺c-kit⁺flt3⁺IL-7R⁻ (63). CLP, common lymphoid progenitor; sIgM, surface IgM. (B) BM cells from WT, FL^{-/-}, and FL^{-/-} $\times$ IL-7R α ^{-/-} mice were investigated for the presence of B cell progenitors as described previously (29, 50); pre-pro-B cells (B220⁺CD43⁺AA4.1⁺CD19⁻), pro-B cells (B220⁺CD43⁺AA4.1⁺CD19⁺), pre-B cells (B220⁺CD43⁻CD19⁺), and immature/mature B cells (B220⁺sIgM⁺). Results are mean values (SD) from six to twelve mice for each group and are expressed as percentages of WT controls (no significant differences in BM cellularity between different genotypes). *Indicates statistically significant differences ($P < 0.01$) between FL^{-/-} and FL^{-/-} $\times$ IL-7R α ^{-/-} mice. 0, no pro-B, pre-B, or B220⁺IgM⁺ cells were detected in FL^{-/-} $\times$ IL-7R α ^{-/-} mice.

these mice. Thus, we next examined the levels of production of IgM, IgG, and IgA in the serum of either FL^{-/-}, IL-7R α ^{-/-}, or FL^{-/-} $\times$ IL-7R α ^{-/-} mice by a sensitive ELISA assay. In contrast to FL^{-/-} or IL-7R α ^{-/-} mice, which produce normal levels of serum IgM and IgG, absolutely no IgM or IgG was detectable in the serum of FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Table III). Similarly, IgA, which was present at slightly reduced levels in IL-7R α ^{-/-} mice, was undetectable in FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Table III). Similar findings were observed for IgD (unpublished data), further confirming the complete absence of mature functional B cells in adult FL^{-/-} $\times$ IL-7R α ^{-/-} mice.

Absence of Detectable Committed B Cell Progenitors in FL^{-/-} $\times$ IL-7R α ^{-/-} Adult BM and Fetal Liver. Since no mature B cells were detected in adult FL^{-/-} $\times$ IL-7R α ^{-/-} mice, we next investigated whether committed B cell progenitors could be detected in FL^{-/-} $\times$ IL-7R α ^{-/-} BM. Recent studies have demonstrated that little or no B cell development takes place in the BM of IL-7^{-/-} and IL-7R α ^{-/-} mice after 7 wk of age, reflected in virtually no detectable B cell progenitors (28, 29). In agreement with this, using a similar staging (29, 49, 50) we found that B220⁺CD43⁺AA4.1⁺CD19⁻ pre-pro-B cells were re-

duced by >99% in FL^{-/-} $\times$ IL-7R α ^{-/-} BM, whereas B220⁺CD43⁺AA4.1⁺CD19⁺ pro-B and pre-B cells were nondetectable (Fig. 4 B). Interestingly, FL^{-/-} mice also showed a dramatic (97.5%) reduction in B220⁺CD43⁺AA4.1⁺CD19⁻ pre-pro-B cells and, as previously shown (37), also reduced levels of pro-B (76%) and pre-B (62%) cells (Fig. 4 B).

Compared with FL^{-/-} and IL-7R α ^{-/-} mice, only very low levels of D-J rearrangement could be detected in the BM of FL^{-/-} $\times$ IL-7R α ^{-/-} mice (Fig. 5 A). The development of mature B cells is dependent on the expression of several transcription factors (25, 51). The B cell lineage-specific activator protein encoded by the Pax5 gene (52) appears to be crucial for B cell development, which in Pax5-deficient mice is blocked at the pro-B cell stage (53, 54). Pax5 RNA was highly expressed in the BM of FL^{-/-} mice, reduced in IL-7R α ^{-/-} mice, and was completely undetectable in FL^{-/-} $\times$ IL-7R α ^{-/-} BM, in agreement with the complete absence of pro-B cells and all subsequent stages of B cell development (Fig. 5 B). Similarly, mb-1 and CD19, two direct target genes of Pax5 (55–58), were expressed in FL^{-/-} BM cells, reduced in IL-7R α ^{-/-} BM cells, but completely absent in FL^{-/-} $\times$ IL-7R α ^{-/-} mice

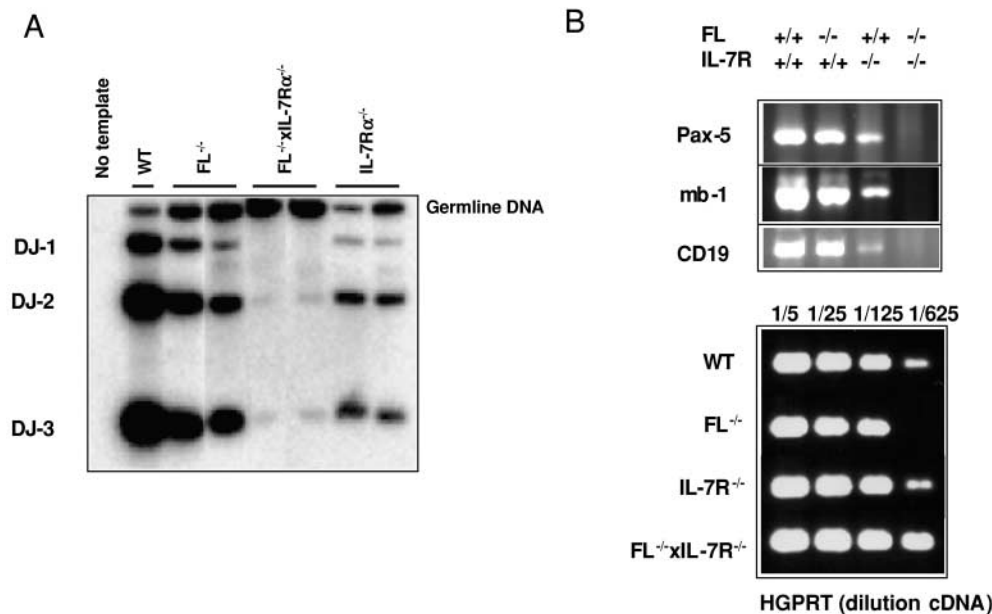


Figure 5. Absence of expression of Pax5 and Pax5 target genes in BM of adult FL^{-/-} × IL-7Rα^{-/-} mice. (A) DNA was isolated from BM of WT, FL^{-/-}, IL-7Rα^{-/-}, and FL^{-/-} × IL-7Rα^{-/-} mice and D-J rearrangement assessed by PCR (described in Materials and Methods). Representative results from two out of seven analyzed mice. (B) RNA was isolated from BM of WT, FL^{-/-}, IL-7Rα^{-/-}, and FL^{-/-} × IL-7Rα^{-/-} mice (described in Material and Methods) and expression of Pax5, mb-1, and CD19 (top) or HGPRT (bottom) analyzed by RT-PCR. Data are from one of three experiments with similar results.

(Fig. 5 B). Thus, flt3 and IL-7Rα signaling appears to be essential for commitment to a Pax5-dependent B cell stage.

Due to the complete absence of not only conventional B cells but also B1 cells in adult FL^{-/-} × IL-7Rα^{-/-} mice, we next compared fetal B lymphopoiesis in IL-7Rα^{-/-} and FL^{-/-} × IL-7Rα^{-/-} mice. B220⁺ IgM⁺ B cells were present, although at reduced levels, in the fetal spleen and liver of IL-7Rα^{-/-} mice, in striking contrast to FL^{-/-} × IL-7Rα^{-/-} mice in which no B220⁺ IgM⁺ B cells were detectable (Fig. 6 A). Furthermore, in contrast to IL-7Rα^{-/-}-deficient mice, FL^{-/-} × IL-7Rα^{-/-} fetal livers lacked detectable B220⁺ CD19⁺ and B220⁺ AA4.1⁺ cells, suggesting that generation of committed B cell progenitors is strictly dependent on IL-7Rα and flt3 signaling also during fetal development (Fig. 6 B).

Discussion

Several cytokines have been shown to promote B cell development in vitro and in vivo (25, 59), but extensive studies of mice deficient in different cytokine ligands and receptors have so far failed to support an indispensable role for cytokines in B cell development. Although the production of mature B cells in the BM of IL-7 and IL-7Rα-deficient mice takes place almost exclusively during fetal and postnatal development (9, 26, 27, 30), a stable pool of mature B cells are maintained throughout adult life in peripheral organs, sustaining normal levels of circulating Ig (9, 26, 27, 30). These B cells might largely be offspring of fetally derived B1 cells, and since present at normal levels in IL-7-deficient mice they have been suggested to represent an important IL-7-independent pathway for B cell development (28). Alternatively, fetal B cell development and maintenance of B cells and Ig production during adult life might be dependent on the concerted action of IL-7 with one or multiple other cytokines. However, mice deficient

in the γc chain receptor, required for signaling of six different lymphokines (including IL-7), express a similar phenotype to that of IL-7Rα^{-/-} mice (31, 32), suggesting that among known lymphokines only IL-7 plays a nonredundant function in B cell development. As the result of these observations, an alternative hypothesis emerged, namely that signaling through cytokine receptors might not be strictly required for the development of mature B cells. Through similar findings, the fundamental importance of cytokine signaling has also been questioned in the development of a number of other blood cell lineages (6–8, 10–12, 14–17, 20–24).

In the present study, we unequivocally demonstrate that signaling through two separate families of cytokine receptors (i.e., an “early acting” receptor tyrosine kinase and a “later acting” hematopoietin receptor) are indispensable for fetal and adult B cell development. This conclusion is based on multiple lines of evidence obtained using methods capable of detecting minute levels of B cells. In contrast to either FL^{-/-} or IL-7Rα^{-/-} mice, peripheral and mesenteric LNs and Peyer’s Patches were not identifiable in FL^{-/-} × IL-7Rα^{-/-} mice. Furthermore, IgM⁺ B cells were found to be completely lacking in the blood, spleen, and BM of adult FL^{-/-} × IL-7Rα^{-/-} mice, whereas, consistent with previous studies, B cells were present (at reduced levels) at all sites in IL-7Rα^{-/-} mice. Similarly, whereas IgA⁺ plasma cells were found at normal or near normal levels in either FL^{-/-} or IL-7Rα^{-/-} mice, they were undetectable in FL^{-/-} × IL-7Rα^{-/-} mice.

Flt3 and IL-7Rα signaling proved essential not only for development of conventional B cells but also for B1 cells and fetal B lymphopoiesis. B1 cells, arising early during fetal development before the appearance of conventional B cells, are believed to be regulated via distinct pathways to those of conventional B cells (28, 47, 48). In fact, before the present study a crucial role for cytokines in B1 cell genesis had not

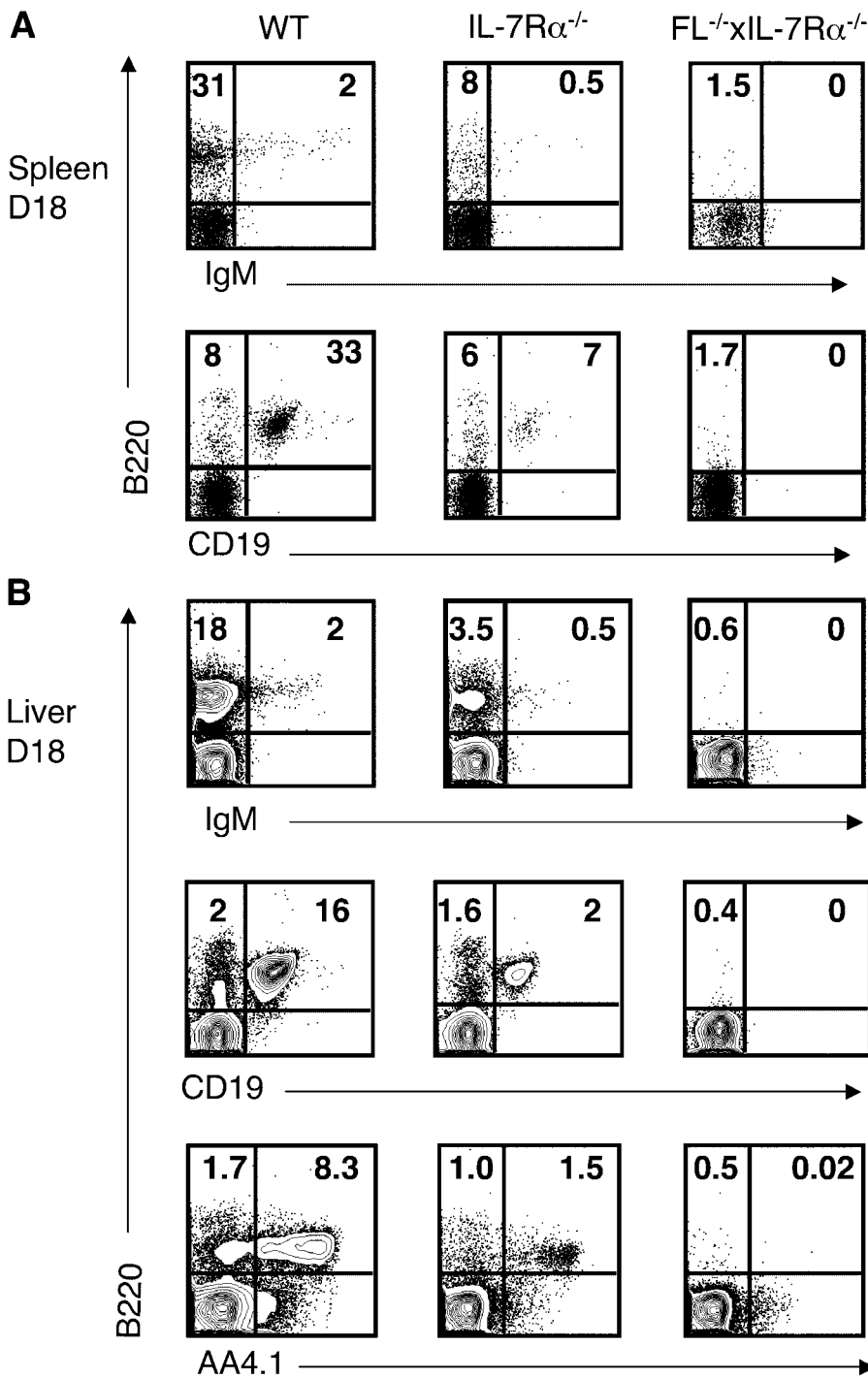


Figure 6. Lack of detectable mature B cells and B cell progenitors during fetal development of mice double deficient in *flt3* ligand and IL-7R α expression. Spleen and liver cells isolated from embryos at day 17–18 of gestation were stained with mAbs against B220, CD19, IgM, and AA4.1 (Materials and Methods). (A) Plots of B220, CD19, and IgM expression in the spleens of WT, IL-7R $\alpha^{-/-}$, and FL $^{-/-}$ $\times$ IL-7R $\alpha^{-/-}$ mice. Numbers represent mean values from three to eight mice. (B) Plots of B220, CD19, and IgM expression in the livers of WT, IL-7R $\alpha^{-/-}$, and FL $^{-/-}$ $\times$ IL-7R $\alpha^{-/-}$ mice. Numbers in A and B represent mean values (percentage of total cells in lymphoid gate) from three to eight mice.

been identified. In this regard, both $\gamma c^{-/-}$ and IL-7-deficient mice have been shown to have normal levels of B1 cells but reduced conventional B cells (31, 28), and herein we show that single FL $^{-/-}$ mice have near normal levels of peritoneal B1 cells, whereas IL-7R $\alpha^{-/-}$ mice have a 63% reduction. In contrast, B1 cells are completely absent in FL $^{-/-}$ $\times$ IL-7R $\alpha^{-/-}$ mice, demonstrating that IL-7R α and *flt3* signaling are indispensable for B1 cell development and/or maintenance. It is noteworthy that in IL-7R $\alpha^{-/-}$ mice we found a significant reduction in B1 cells, whereas B1

cells are normal in IL-7 $^{-/-}$ mice (28). Since another cytokine, thymic stromal lymphopoietin, has been shown to utilize the IL-7R α in its signaling (60), this might suggest that thymic stromal lymphopoietin together with FL and IL-7 might be critically involved in generation of B1 cells.

A critical role for *flt3* and IL-7R signaling in fetal B cell development was further substantiated by a complete absence of mature B220 $^{+}$ IgM $^{+}$ B cells, in striking contrast to IL-7R $\alpha^{-/-}$ mice. Furthermore, whereas the fetal liver of IL-7R $\alpha^{-/-}$ mice contained B220 $^{+}$ CD19 $^{+}$ and B220 $^{+}$

AA4.1⁺ B cell progenitors, no B220⁺ cells coexpressing CD19 or AA4.1 were detected in the fetal liver of FL^{-/-} × IL-7Rα^{-/-} mice, suggesting a critical role of FL and IL-7 in the earliest stages of fetal B cell commitment and development. Finally, and most strikingly, whereas a highly sensitive ELISA assay confirmed that the residual levels of B cells in adult IL-7Rα^{-/-} mice can produce normal Ig levels, no Ig production (IgM, IgG, IgA and IgD) was observed in FL^{-/-} × IL-7Rα^{-/-} mice.

Although certain transcription factors have been demonstrated to be critically involved in B cell commitment and development (25, 51), and our present studies demonstrate that B cell genesis is also strictly dependent on specific cytokine signaling, the potential cross talk between the two remain poorly understood. It is well known, however, that transcription factors can regulate cytokine receptor expression (4), but there is less evidence for a role of cytokine signaling in the regulation of transcription factor expression. It is striking that expression of *Pax5*, the proposed master regulator of B cell commitment and development (53, 54), was completely absent in the BM of FL^{-/-} × IL-7Rα^{-/-} mice, as were the direct *Pax5* target genes *mb-1* and *CD19* (55–58). This could potentially reflect an indispensable role of flt3 and IL-7Rα in regulation of *Pax5* expression, but since pro-B cells and all downstream stages of B cell development were completely lacking in the BM of FL^{-/-} × IL-7Rα^{-/-} mice, we rather favor that flt3 and IL-7Rα are absolutely required to promote B cell development to a *Pax5*-dependent stage. This is interesting in the context that *Pax5* is thought to act as a “B cell lineage locker,” restricting the lineage potentials of otherwise multipotent progenitors to a committed B cell fate (54, 61), implicating that B cell commitment might be dependent on signaling through flt3 and IL-7Rα. Although FL^{-/-} × IL-7Rα^{-/-} BM showed low levels D-J rearrangement, this could potentially derive from cells not committed to the B cell lineage, since D-J rearrangement can be found in T cell progenitors (62).

Despite flt3 and c-kit being coexpressed at early stages of B cell development, including CLP (37, 63), the present findings demonstrate a distinct role of flt3 but not c-kit in early B cell development, since γc^{-/-} × c-kit^{-/-} mice produce B cells at levels comparable to that of single γc^{-/-} mice (64). Although the mechanisms by which flt3 and IL-7Rα interact to maintain B cell production remains to be established, the function of the two cytokine receptors are evident at distinct and overlapping stages of B cell development (34, 65). Flt3 is expressed (34, 63) at earlier stages than IL-7Rα, and FL-deficient mice have a defect primarily in early B cell progenitors, including the CLP (37). In contrast, IL-7Rα is primarily expressed at later stages of B cell development, and IL-7-deficient mice have normal levels of CLPs (29) but reduced levels of B cell progenitors and mature B cells (26, 27). However, the complete absence of conventional and B1 cells in fetal and adult FL^{-/-} × IL-7Rα^{-/-} mice most likely also reflects important synergistic interactions and potential cross-talk between these two cytokine receptor signaling pathways during multiple stages of B cell development (from CLP to pre-B cells) where both

receptors are coexpressed. This is supported by the potent in vitro synergy between FL and IL-7 in promoting B cell development and expansion from adult uncommitted BM progenitors in which the combined action of FL and IL-7 proved both sufficient and absolutely essential (39).

The potent synergy between one of the cytokine tyrosine kinase receptors flt3 and/or c-kit and multiple members of the hematopoietic receptor superfamily in promoting growth and development of different blood cell lineages is a well-described in vitro phenomenon (2, 3, 34). The present findings propose an important physiological synergy between an early acting tyrosine kinase receptor and late acting hematopoietin receptor during B cell development, rather than between intensively explored combinations of hematopoietic lineage factors. This apparent synergy may not only be critical for B cell development but may also prove essential for the development of other blood cell lineages. In support of this hypothesis, a combined deficiency in both c-kit and γc chain receptor expression has been shown to result in a profound block in early thymocyte development (64). Although c-kit^{-/-} mice die 2–6 d after birth (due to anemia) complicating further analysis and it remains unclear which (one or multiple) of the 6 cytokines acting through the γc chain receptor are involved in thymocyte development, this finding lends further support toward an essential role of interactions between early acting cytokine tyrosine kinase receptors and later acting hematopoietin receptors in blood lineage development.

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