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B cells require type 1 interferon to produce alloantibodies to transfused KEL-expressing RBCs in mice

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Abstract

**Background**—Alloantibodies to RBC antigens can cause significant hemolytic events. Prior studies have demonstrated that inflammatory stimuli in animal models and inflammatory states in humans, including autoimmunity and viremia, promote alloimmunization. However, molecular mechanisms underlying these findings are poorly understood. Given that type 1 interferons (IFNα/β) regulate antiviral immunity and autoimmune pathology, the hypothesis that IFNα/β regulates RBC alloimmunization was tested in a murine model.

**Study Design and Methods**—Leuko-reduced murine RBCs expressing the human KEL glycoprotein were transfused into control mice (WT), mice lacking the unique IFNα/β receptor (IFNAR1−/−), or bone marrow chimeric mice lacking IFNAR1 on specific cell populations. Anti-KEL IgG production, expressed as mean fluorescence intensity (MFI), and B cell differentiation were examined.

**Results**—Transfused WT mice produced anti-KEL IgG alloantibodies (peak response MFI=50.4). However, the alloimmune response of IFNAR1−/− mice was almost completely abrogated (MFI=4.2, p<0.05). The response of bone marrow chimeric mice lacking IFNAR1 expression in all hematopoietic cells or specifically in B cells was also diminished (MFI=3.8 and 5.4, respectively, compared to control chimeras, MFI=65.6, p<0.01). Accordingly, transfusion-
induced differentiation of IFNAR1−/− B cells into germinal center B cells and plasma cells was significantly reduced, compared to WT B cells.

Conclusions—This study demonstrates that B cells require signaling from IFNα/β to produce alloantibodies to the human KEL glycoprotein in mice. These findings provide a potential mechanistic basis for inflammation-induced alloimmunization. If these findings extend to human studies, patients with IFNα/β-associated conditions may have an elevated risk of alloimmunization and benefit from personalized transfusion protocols.

Keywords
Type I interferon; alloimmunization; KEL; inflammation

Introduction
With approximately 15 million units of red blood cells (RBC) transfused per year, RBC transfusion is the most common procedure performed in United States hospitals. Unlike ABO and Rh antigens, other common RBC antigens, including Kell, Duffy, and Kidd, are not routinely matched between donor units and recipients. Exposure to allogeneic RBC antigens during transfusion leads to production of antigen-specific alloantibodies in 3–10% of transfusion recipients and 30–50% of patients with sickle-cell disease. Subsequent exposure to the RBC antigen may result in potentially fatal hemolytic transfusion reactions. Additionally, transfusion-dependent patients with hemoglobinopathies, including sickle cell disease, may have rare or multiple RBC alloantibodies. Thus, compatible RBC units for these patients can be exceedingly limited or unattainable, resulting in anemia-associated morbidity and mortality. Elucidating factors that influence alloantibody production could improve prediction of these alloimmune responses, and allow for intervention to reduce alloimmunization and hemolytic events.

One such factor is the inflammatory state of the transfusion recipient. Ramsey and Smietana reported an increased prevalence of autoimmune disease in alloimmunized patients. More recent studies have reported an increased incidence of alloimmunization in patients with inflammatory bowel disease, febrile transfusion reactions, and sickle cell-mediated acute chest syndrome. Recently, Evers et al. reported that patients with viremia also have an elevated risk of alloimmunization. However, patients with Gram-negative bacteremia had significantly lower rates of alloimmunization. Accordingly, studies in mouse models of alloimmunization have demonstrated that viral mimetics promote RBC alloimmune responses, while lipopolysaccharide (LPS) inhibits responses. These findings suggest that specific inflammatory pathways associated with certain conditions, including some autoimmune diseases and viral infection, promote RBC alloimmune responses. Recently, Arneja et al demonstrated a role for interleukin (IL)-6 in T cell responses to stored RBCs. However, other cytokine-mediated mechanisms underlying inflammation-induced alloimmunization have not been investigated.

The inflammatory conditions that promote alloimmunization are associated with the production of numerous inflammatory cytokines, including tumor necrosis factor α (TNFα), IL-12, and IL-6. One class of inflammatory cytokines that regulates both antiviral immunity
and autoimmune pathology is the type I Interferon (IFNα/β) family, consisting of 13 IFNα genes and 1 IFNβ gene. In a virally infected cell, recognition of viral nucleic acids by ubiquitously expressed pattern recognition receptors (PRRs), including Toll-like receptors, results in production and secretion of IFNα/β. Binding of IFNα/β to its dimeric receptor, consisting of IFNα Receptor (IFNAR) 1 and IFNAR2, induces expression of interferon stimulated genes (ISGs) that inhibit viral replication and dissemination. Long-term viral protection is dependent on adaptive immune responses that produce neutralizing antibodies. Studies in murine models of viral infection and immunization have shown that IFNα/β can enhance or inhibit antibody responses. Multiple studies have reported that injection of IFNα/β promotes antibody responses to some soluble antigens by increasing activation of dendritic cells and lymphocytes. However, Muller et al. generated IFNAR1-deficient mice (IFNAR1−/−) and reported that wildtype (WT) and IFNAR1−/− mice produced comparable IgM and IgG responses to immunization with multiple soluble antigens, including ovalbumin and a viral glycoprotein. More recent studies have demonstrated that IFNAR1−/− mice produce elevated levels of protective antibodies in chronic viral infections. Thus, IFNAR1−/− mice can mount antibody responses to multiple viral and soluble antigens.

IFNAR signaling has also been implicated in the pathogenesis of multiple autoimmune diseases, including myositis, scleroderma, rheumatoid arthritis, and Sjogren’s syndrome. Patients with systemic lupus erythematosus (SLE) frequently have elevated serum IFNα/β and ISG expression, which correlate with increased auto-antibody levels and disease severity. This association has led to the use of anti-IFNα/β therapy in clinical trials.

The role of IFNα/β in RBC alloimmunization has not been previously investigated. In comparison to viral and soluble antigens, RBC-bound antigens differ in their route of administration, quantity of antigen, duration of exposure, and accessibility to niches within secondary lymphoid tissue. Thus, the role of IFNα/β in RBC alloimmunization may differ from their role in autoimmunity, viral infections, or immunization. Given that viral infections and multiple autoimmune diseases induce IFNα/β production and promote RBC alloimmunization, we tested the hypothesis that IFNα/β regulates alloimmune responses to RBC antigens. Given the large number of antigenic differences between human donors and recipients, we utilized a murine transfusion model that allows examination of alloimmune responses to a single donor RBC antigen. Using this model, we assessed the role of IFNα/β in promoting RBC alloimmunization. Further, we assessed the contribution of cell-specific IFNα/β signaling to the RBC alloimmune response.

Materials and Methods

Mice

C57BL/6 and congenic C57BL/6-Ly5.1 WT mice were purchased from Charles River Laboratories (Wilmington, MA). muMT−, TCRα−/−, and Zbtb46-DTR mice were purchased from Jackson Laboratories (Bar Harbor, ME). IFNAR1−/− and KEL RBC transgenic mice (previously published as KEL2B mice) were previously described. All mice had been...
backcrossed to the C57BL/6 background for more than 8 generations. All animal protocols were approved by the Yale Institutional Animal Care and Use Committee (IACUC). In accordance to the IACUC protocol, experimental groups consisted of 4–5 mice per independent experiment.

**RBC Transfusion**

Peripheral blood of KEL RBC mice was collected in 12% Citrate Phosphate Dextrose Adenine (CPDA-1, Jorgensen Labs, Melville, NY) by retro-orbital bleeding and leuko-reduced with a Pall (East Hills, NY) syringe filter. WT, IFNAR1\(^{-/-}\), or chimeric mice were transfused i.v. with 75 μL leuko-reduced packed RBCs, the mouse equivalent of 1 unit of human RBCs. For secondary responses, mice were re-transfused 28 days later. For IFNAR1 blocking experiments, WT mice were i.p. injected with 600 μg anti-mouse IFNAR1 (clone: MAR1-5A3, BioXCell, West Lebanon, NH), isotype control mouse IgG1 (MOPC-21, BioXCell) or PBS 24 hours prior to and 3 and 7 days after transfusion.

**Generation of bone marrow chimeras**

Recipient WT C57BL/6 (CD45.2\(^{+}\)) or IFNAR1\(^{-/-}\) (CD45.2\(^{+}\)) mice were exposed to 2 rounds of X-ray irradiation (6.35 Gy, 3 hours apart) using an X-RAD 320 irradiator (Precision X-ray Inc., North Branford, CT). Recipients were i.v. injected 2–4 hrs after irradiation with a total of 3×10\(^{6}\) bone marrow cells from WT C57BL/6-Ly5.1 (CD45.1\(^{+}\)), IFNAR1\(^{-/-}\), muMT\(^{-}\) (CD45.2\(^{+}\)), TCR\(^{\alpha^{-/-}}\) (CD45.2\(^{+}\)) and/or Zbtb46-DTR (CD45.1\(^{+}\), CD45.2\(^{+}\)) mice. Peripheral blood was analyzed for lymphocyte reconstitution 6 weeks after bone marrow transfer. Mice were transfused 8–9 weeks following bone marrow reconstitution. For chimeras generated with Zbtb46-DTR bone marrow, DTR-expressing cells were depleted with 2 i.p. injections of diphtheria toxin (DT, Sigma-Aldrich, St. Louis, MO) 3 (60 ng/g) and 1 day (40ng/g) prior to transfusion as previously described\(^{39,40}\). Doses of DT were titrated in Zbtb46-DTR chimeras to achieve efficient conventional DC depletion.

**Measurement of anti-KEL alloantibodies**

Transfusion of murine KEL RBCs induces anti-KEL antibody production in allogeneic mice that do not express the human KEL glycoprotein\(^{38}\). These anti-KEL antibodies formed following transfusion of KEL RBCs are defined as anti-KEL alloantibodies. Serum anti-KEL IgG was measured by flow cytometric crossmatch 7, 14, 21, and 28 days after transfusion as previously described\(^{41}\). Briefly, serum from transfused mice was incubated with KEL RBCs and subsequently stained for RBC-bound IgG (Goat anti-mouse IgG APC, Jackson ImmunoResearch, West Grove, PA). Undiluted serum was used to maximize the sensitivity of the assay. The adjusted MFI was calculated by subtracting the reactivity of serum with syngeneic C57BL/6 RBCs from the reactivity of serum with KEL RBCs. Figure data illustrates the peak antibody response, 21–28 days following transfusion. Flow cytometry of RBCs was performed using a BD FACS Calibur (San Jose, CA) and analyzed using FlowJo software (Tree Star, Ashland, OR).
Flow cytometric Analysis

Single cell suspensions of peripheral blood leukocytes and splenocytes were analyzed following RBC lysis. Bone marrow was collected by flushing tibias with media. For dendritic cell analysis, spleens were minced with a razor blade prior to cell filtration with a 70 μM nylon mesh. Suspensions were stained with fluorescently conjugated antibodies specific for cell surface proteins, including CD45.2 (Clone: 104), CD138 (Syndecan-1), CD19 (6D5), TCRβ (H57-597), IgD (11-26c.2a), I-A/I-E (MHC II, M5/114.15.2), and GL7 from Biolegend (San Diego, CA); CD45.1 (A20), CD11c (N418), B220 (RA3-6B2), and IFNAR1 (MAR1-5A3) from Ebioscience (San Diego, CA), and Fas/CD95 (Jo2) from BD Biosciences (San Jose, CA). Zombie-NIR (Biolegend) was used to exclude dead cells. Cells were acquired with a Miltenyi MACSQuant flow cytometer and analyzed using FlowJo.

In vitro B cell cultures

B cells were isolated from WT and IFNAR1−/− spleens by negative selection using a magnetic cell isolation kit (StemCell Technologies, Vancouver, BC). B cell purity analyzed by flow cytometry was 90–95%. B cells (1×10⁶/mL) were cultured in complete RPMI previously described containing anti-CD40 (10 μg/mL FGK4.5) with or without recombinant IFNα (100 U/mL, HC1040, Hycult Biotech, Netherlands) for 72 hrs. Development of plasma cells (CD19+IgDloB220loCD138+) was measured by flow cytometry.

Statistics

Statistical analysis was performed using GraphPad Prism software (San Diego, CA). Statistical significance between two groups was determined using an unpaired t-test or Mann Whitney U test for parametric and non-parametric data, respectively. Significance between three groups was determined using a Kruskal-Wallis test with a Dunn’s post-test. All anti-KEL IgG, germinal center B cell, and primary plasma cell data was analyzed using non-parametric tests. B cell culture data was analyzed with an unpaired two-tailed t-test. Bars on graphs indicate the mean of the data, and the circles represent individual data points from one mouse or one culture. Statistical significance was determined for all individually performed experiments containing at least 4 mice per group. With the exception of B cell culture data, Figures in the primary text display results of one experiment of 2–3 completed experiments with 3–5 mice per group. Results of repeated experiments are shown in Supplemental figures.

Results

IFNα/β regulates alloimmunization to KEL-expressing RBCs

IFNα/β is induced in inflammatory conditions associated with increased alloimmunization. Thus, we examined whether expression of the only receptor for IFNα/β, IFNAR, was required for RBC alloimmunization. We utilized a murine model in which donor murine RBCs express the human KEL glycoprotein in a RBC specific manner (KEL RBCs) 38. As shown in Figure 1A, WT mice transfused with leuko-reduced KEL RBCs produced anti-KEL glycoprotein IgG alloantibodies in the absence of an adjuvant. In contrast, in 3 out of 3
experiments, the anti-KEL antibody response of IFNAR1−/− mice was almost completely abrogated after transfusion of KEL RBCs (Figure 1A and Supplemental Figure 1A). The abrogated response could be due to altered lymphoid structure or leukocyte development in IFNAR1−/− mice. To address this possibility, the anti-KEL IgG response was examined in WT mice treated with an IFNAR1 blocking antibody (MAR1-5A3), an isotype control IgG1 antibody, or PBS. Compared to control groups, MAR1-5A3 treated mice produced significantly lower levels of anti-KEL IgG following transfusion (Figure 1B and Supplemental Figure 1B). Collectively, these results indicate that IFNα/β critically regulates alloimmunization to KEL-expressing RBCs.

**Hematopoietic cell expression of IFNα/β receptors in transfusion recipients is required for KEL RBC alloimmunization**

IFNAR1 is expressed by multiple hematopoietic and non-hematopoietic cells. Additionally, IFNAR signaling has been reported to promote humoral immune responses by activating multiple cell types. To determine whether hematopoietic or non-hematopoietic IFNAR1 expression is required for alloimmunization following KEL RBC transfusion, alloimmune responses in bone marrow chimeras were examined. Recipients were irradiated and reconstituted with donor bone marrow 8 weeks prior to transfusion. As shown in Figure 1C, reconstitution of irradiated IFNAR1−/− mice with WT bone marrow rescued the alloimmune response of IFNAR1−/− mice. In contrast, irradiated WT mice reconstituted with IFNAR1−/− bone marrow failed to produce anti-KEL IgG following transfusion (Figure 1C and Supplemental Figure 1C). Thus, IFNAR1 expression by hematopoietic cells is required for KEL-RBC alloimmunization.

**IFNAR expression in conventional dendritic cells is dispensable for RBC alloimmunization**

To determine which hematopoietic cell type(s) utilize(s) IFNα/β to promote alloantibody production, responses were assessed in mixed bone marrow chimeras that lack IFNAR1 expression in specific cell types. Given that prior studies have established a role for IFNα/β in promoting dendritic cell maturation and antigen presentation, we hypothesized that IFNAR signaling in conventional dendritic cells (cDCs) may regulate alloimmunization. To generate chimeras lacking IFNAR1 expression in cDCs, we utilized Zbtb46-DTR mice, which specifically express the diphtheria toxin receptor (DTR) on cDCs. WT mice were irradiated and reconstituted with a mixture of Zbtb46-DTR (CD45.1+, CD45.2+) bone marrow and either WT (CD45.1+) or IFNAR1−/− (CD45.2+) bone marrow. CD45 congenic markers were used to identify cells derived from different donors (Figure 2A). After 8 weeks of reconstitution, analysis of IFNAR1 expression on splenocytes from Zbtb46-DTR x IFNAR1−/− chimeras confirmed that CD45.2+ cells were derived from IFNAR1−/− BM (Figure 2B). Zbtb46-DTR-expressing cDCs were then depleted by injecting diphtheria toxin (DT) prior to transfusion. As a result, nearly all remaining cDCs in DT-treated Zbtb46-DTR x IFNAR1−/− chimeras were derived from IFNAR1−/− BM (Figure 2C). However, following transfusion, DT and PBS-treated chimeras produced comparable levels of anti-KEL IgG. The anti-KEL IgG levels were also comparable to those in DT-treated chimeras containing WT cDCs (Zbtb46-DTR x WT, Figure 2D and Supplemental Figure 2A). These results indicate that IFNAR1 expression on cDCs is not required for alloantibody responses to KEL-expressing RBCs.
T cell IFNAR expression is not required for alloimmunization to KEL RBCs

Studies have demonstrated that IFNAR signaling in T cells can regulate humoral immune responses in viral infection and immunization models \(^{21,46}\). To determine whether T cell IFNAR signaling regulates alloimmunization to KEL RBCs, mixed chimeras that lack IFNAR1 expression specifically on T cells were generated. Irradiated IFNAR1\(^{-/-}\) (CD45.2\(^+\)) recipients were reconstituted with bone marrow from T cell-deficient TCR\(^{\alpha^{-/-}}\) (CD45.2\(^+\)) mice and either WT (CD45.1\(^+\)) or IFNAR1\(^{-/-}\) bone marrow. IFNAR1\(^{-/-}\), rather than WT mice were used as bone marrow recipients to ensure that irradiation-resistant recipient thymocytes in TCR\(^{\alpha^{-/-}}\) \times IFNAR1\(^{-/-}\) chimeras lacked IFNAR1 expression. Following reconstitution, T cells in TCR\(^{\alpha^{-/-}}\) \times WT chimeras were predominately derived from donor CD45.1\(^+\) WT bone marrow. However, recipient CD45.2\(^+\) irradiation-resistant T cells were also present (Figure 3A). TCR\(^{\alpha^{-/-}}\) \times IFNAR1\(^{-/-}\) chimeras were only generated with CD45.2\(^+\) bone marrow (Figure 3B). However, measurement of IFNAR1 expression in T cells confirmed that TCR\(^{\alpha^{-/-}}\) \times IFNAR1\(^{-/-}\) chimeras contained only IFNAR1-deficient T cells. (Figure 3C). Following transfusion of KEL RBCs, both groups of chimeras produced comparable amounts of anti-KEL IgG (Figure 3D and Supplemental Figure 2B). These results indicate that IFNAR expression by T cells is not required for alloimmunization to KEL RBCs.

IFNAR expression by B cells is required for alloimmunization to KEL RBCs

Studies of viral infection and immunization have also shown that IFN\(\alpha/\beta\) can directly promote B cell activation and antibody production \(^{21,47,48}\). To examine whether IFNAR signaling in B cells regulates RBC alloimmunization, mixed chimeras were generated by reconstituting irradiated CD45.2\(^+\) WT mice with bone marrow from B cell-deficient muMT\(^-\) (CD45.2\(^+\)) mice and either WT (CD45.1\(^+\)) or IFNAR1\(^{-/-}\) (CD45.2\(^+\)) mice. Chimeras reconstituted with only IFNAR1\(^{-/-}\) bone marrow were also generated to serve as controls. Following reconstitution, nearly all CD19\(^+\) B cells in muMT\(^-\) \times WT chimeras were derived from CD45.1\(^+\) WT bone marrow (Figure 4A). In muMT\(^-\) \times IFNAR1\(^{-/-}\) chimeras, expression of CD45.2 on all cells excluded a similar analysis (Figure 4B). However, analysis of IFNAR1 expression demonstrated that all B cells in muMT\(^-\) \times IFNAR1\(^{-/-}\) chimeras were IFNAR1-deficient (Figure 4C). Upon transfusion with KEL RBCs, muMT\(^-\) \times WT chimeras produced alloantibodies. However, like IFNAR1\(^{-/-}\) control chimeras, muMT\(^-\) \times IFNAR1\(^{-/-}\) chimeras produced significantly lower levels of anti-KEL IgG (Figure 4D and Supplemental Figure 2C). Thus, IFNAR signaling in B cells regulates alloimmunization to KEL RBCs.

IFN\(\alpha/\beta\) promotes germinal center B cell and plasma cell differentiation

Upon antigen recognition, follicular B cells differentiate into germinal center (GC) B cells that ultimately produce antibody producing plasma cells. To determine whether IFN\(\alpha/\beta\) regulates B cell differentiation, we examined GC B cell and plasma cell formation in IFNAR1\(^{-/-}\) and WT mice following KEL RBC transfusion. Prior to transfusion, similar levels of GC B cells (CD19\(^+\)IgD\(^{lo}\)Fas\(^{-hi}\)GL7\(^{hi}\)) were detected in spleens of IFNAR1\(^{-/-}\) and WT mice (Figure 5A–B and Supplemental Figure 3A). Compared to WT controls, the percentage of GC B cells in IFNAR1\(^{-/-}\) mice was significantly decreased, 8 days following transfusion. The total number of GC B cells was statistically different between the 2 groups.
in 1 of 2 experiments (Figure 5C–D and Supplemental Figure 3B). To enrich for KEL-specific germinal center B cells and plasma cells, mice were re-transfused with KEL RBCs on Day 28. Eight days following a second transfusion, the percentage and number of GC B cells in IFNAR1−/− mice were reduced compared to WT controls (Figure 5E,F and Supplemental Figure 3C). In addition, despite a potential increase in IFNAR1−/− GC B cells following a second transfusion, anti-KEL IgG responses by IFNAR1−/− mice remained fully abrogated (Figure 5G and Supplemental Figure 3D). Differentiation of plasma cells was also assessed 14 days after the second transfusion. The percent and total number of bone marrow plasma cells (CD19+IgDloB220loCD138+) were significantly reduced in IFNAR1−/− mice, compared to WT controls (Figure 6A–C and Supplemental Figure 4). These results indicate that IFNα/β regulates KEL alloimmune responses by promoting GC B cell and plasma cell differentiation.

Given that IFNAR expression is abrogated in all cell types of IFNAR1−/− mice, it is possible that IFNAR expression in other hematopoietic and non-hematopoietic cells may influence plasma cell differentiation. To address this possibility, we isolated WT and IFNAR1−/− spleen B cells and examined plasma cell differentiation in ex vivo cultures in the presence and absence of recombinant IFNα (rIFNα). Magnetically selected B cells were cultured for 72 hrs in the presence of the anti-CD40 antibody, FGK4.5, to promote cell survival. In accordance with prior studies 48,49, the addition of rIFNα to WT cultures resulted in elevated production of plasma cells (CD19+IgDloB220loCD138+), compared to cultures lacking rIFNα. However, the addition of rIFNα did not increase plasma cell development in IFNAR1−/− B cell cultures (Figure 6D–F). This result demonstrates that IFNα/β directly promotes B cell differentiation into antibody-producing plasma cells.

**Discussion**

Identifying patients with an elevated risk of transfusion would allow interventions, such as extended antigen matching, to inhibit alloimmunization and hemolytic events. However, diagnostic tests to predict alloimmunization have not been developed. This is in part due to the lack of understanding of cellular and molecular pathways that promote alloimmunization. In this study, we demonstrate that recipient expression of interferon receptors (IFNAR) is required for alloimmunization to the human KEL glycoprotein in a murine transfusion model. Although the receptor for IFNα/β is expressed by many non-hematopoietic and hematopoietic cell types, we demonstrate that IFNAR expression by B cells critically regulates the humoral alloimmune response. We further show that IFNAR promotes germinal center B cell and plasma cell differentiation following transfusion.

IFNα/β has been shown to have diverse effects on humoral immune responses to varying infectious organisms and immunogenic antigens 21–24. Given that IFNAR1−/− and WT mice were reported to produce similar antibody responses in many other models 22, the abrogated RBC alloimmune response of IFNAR1−/− mice was unlikely due to potentially altered lymphoid architecture or hematopoiesis in IFNAR1−/− mice. Rather, our interpretation of these data is that binding of IFNα/β to IFNAR activates downstream signaling that is required for alloimmunization to KEL RBCs. This conclusion is supported by the finding...
that treatment of WT mice with an IFNAR1 blocking antibody significantly inhibited the anti-KEL IgG response.

These findings provide insight into previously reported studies in mouse transfusion models. Treatment of recipient mice with inflammatory pathogen associated molecular patterns (PAMPs), including poly(I:C) and CpG, promotes alloimmunization to RBCs expressing KEL or other alloantigens. Poly(I:C) is a mimetic of viral double stranded RNA (dsRNA) that induces robust production of IFNα/β by many cell types. Our demonstration that IFNAR expression is required for KEL RBC alloimmunization raises the possibility that poly(I:C) promotes alloimmunization by inducing IFNα/β. However, this should be formally tested in transfusion models that require the use of poly(I:C) to induce alloimmunization.

Multiple studies have successfully utilized mixed bone marrow chimeras to examine the role of IFNAR signaling in specific cell types. Using this approach, we found that while IFNAR expression by B cells was critical for anti-KEL alloimmune responses, IFNα/β-mediated responses by cDCs and T cells were dispensable. In contrast, prior studies utilizing IFNα/β injections to increase antibody responses have reported that IFNα/β-mediated responses by cDCs, T cells, and B cells promoted humoral immune responses to soluble antigens. This apparent discrepancy might reflect inherent differences between responses to soluble and RBC-bound antigens. In addition, although RBC alloimmunization to other antigens requires DC presentation to T cells, it is possible that T cell help is not required for anti-KEL responses in this model. Further, IFNAR signaling in T cells or DCs may be able to promote B cell alloantibody production. In this case, deletion of IFNAR from either cell type would not inhibit alloimmunization. These findings may also indicate that IFNα/β-mediated contributions of T cells and cDCs to the alloimmune response may depend on the level of available IFNα/β, which may be produced and consumed in localized niches accessible to B cells. Unlike IFNα/β treated mice, we observed that the level of serum IFNα in mice transfused with KEL RBCs was nearly undetectable (unpublished results). However, our IFNAR1−/− results indicate that there is a source of IFNα/β that stimulate B cells to produce anti-KEL antibodies. We hypothesize that transfused RBCs might be associated with damage associated molecular patterns (DAMPs) that can bind to PRRs and induce IFNα/β production. Such DAMPs may be present on RBCs, platelets, or remaining leukocytes. Future studies are needed to identify the sensor(s) and ligand(s) that promote IFNα/β production in alloimmunization.

Analysis of B cell differentiation in IFNAR1−/− mice revealed that IFNα/β regulates GC B cell and plasma cell differentiation in response to KEL RBC transfusion. Additionally, in accordance with prior studies of human B cells co-cultured with IFNα/β-producing plasmacytoid DCs, rIFNα directly induced plasma cell differentiation in cultures of isolated spleen B cells. These findings are consistent with results from viral infection and immunization models and provide insight into the role of IFNα/β in transfusion-induced B cell differentiation. We hypothesize that IFNα/β may regulate the selection, proliferation and/or maintenance of KEL-specific GC B cell clones. However, plasma cell formation is the result of numerous B cell processes including antibody class switching, affinity maturation, and cell migration. Thus, other mechanisms underlying IFNα/β-induced
B cell differentiation during alloimmunization are plausible and are the focus of further study.

In summary, we demonstrate that expression of the IFNα/β receptor by B cells is required for alloimmunization to the human KEL glycoprotein on transfused murine RBCs. We further show that IFNα/β promotes GC B cell and plasma cell differentiation following RBC transfusion. These findings provide a potential mechanistic basis for inflammation-induced alloimmunization. If they extend to human studies, patients with IFNα/β-associated conditions, including viral infection and autoimmunity, or those treated with IFN-based therapies, may have an elevated risk of alloimmunization.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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References


Figure 1. IFNα/β promotes alloimmunization to KEL RBCs
Peripheral blood of KEL-expressing transgenic mice was leuko-reduced and transfused into recipients on Day 0. Serum anti-KEL IgG was measured by flow cytometric crossmatch. The adjusted MFI was calculated by subtracting the reactivity of serum with WT RBCs from serum reactivity with KEL RBCs. (A) Anti-KEL IgG in serum of WT or IFNAR1−/− mice 28 days after transfusion with KEL RBCs. (B) Serum anti-KEL IgG of transfused WT mice injected i.p. with anti-IFNAR1 blocking antibody (MAR1-5A3), an isotype control IgG1 antibody (MOPC-21), or PBS on Day −1, +2, and +7. (C) Anti-KEL IgG of transfused WT, IFNAR1−/−, and bone marrow chimeric mice. Recipients were irradiated and reconstituted with donor bone marrow cells, 8 weeks prior to transfusion. “n/a”: non applicable. One of 2 (B–C) or 3 (A) independent experiments with 4–5 mice/group. Data from repeated experiments are shown in Supplemental Figure 1. Bars on graphs indicate the mean of the data, and the circles represent individual data points from one mouse. *p<0.05 by (A, C) Mann Whitney U test and (B) Kruskal-Wallis test with a Dunn’s post-test.
Figure 2. IFNAR expression by conventional dendritic cells is dispensable for RBC alloimmunization
Mixed chimeras were generated by reconstituting irradiated CD45.2 WT recipients with a 4:1 mixture of Zbtb46-DTR (CD45.1+, CD45.2+) and either IFNAR1−/− (CD45.2+) or WT (CD45.1+) bone marrow. (A) Representative flow cytometric analysis of splenocytes derived from bone marrow donors in Zbtb46-DTR x IFNAR1−/− chimeras. (B) IFNAR1 expression of splenocytes gated in (A). (C) Representative flow cytometric analysis of spleen CD11c+ MHCII+ conventional dendritic cells in Zbtb46-DTR x IFNAR1−/− chimeras treated with PBS or dipheria toxin (DT) 8 weeks after bone marrow transfer. (D) Serum anti-KEL IgG produced by indicated chimeras treated with PBS or DT prior to transfusion with KEL RBCs. Z46-DTR = Zbtb46-DTR. Numbers on plots indicate percent of CD19− TCRβ− cells within the drawn gate. Representative of 3 independent experiments with 4–5 mice/group.
n.s., not significant by Kruskal-Wallis test with a Dunn’s post-test. Data from repeated alloimmunization experiments are shown in Supplemental Figure 2A.
Figure 3. T cell IFNAR expression is not required for RBC alloimmunization

Mixed chimeras were generated by reconstituting irradiated IFNAR1−/− (CD45.2+) recipients with a 4:1 mixture of TCRα−/− (CD45.2+) and either IFNAR1−/− (CD45.2+) or WT (CD45.1+) bone marrow. (A–B) Representative reconstitution analysis of peripheral blood lymphocytes from (A) TCRα−/− x WT and (B) TCRα−/− x IFNAR1−/− chimeras. Numbers on plots indicate percent of live (left), CD45.2+ (middle), or CD45.1+ (right) cells within the drawn gate. (C) Relative expression of IFNAR1 on peripheral blood TCRβ+ T cells from IFNAR1−/− mice and the indicated chimeras. (D) Anti-KEL IgG in serum of indicated chimeras following transfusion with KEL RBCs. Representative of 3 independent experiments with 4–5 mice/group per experiment. n.s., not significant by Mann Whitney U

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Data from repeated alloimmunization experiments are shown in Supplemental Figure 2B.
Figure 4. IFNAR expression by B cells is required for alloimmunization to KEL RBCs
Mixed chimeras were generated by reconstituting irradiated WT (CD45.2+) recipients with a 2:1 mixture of muMT− (CD45.2+) and either IFNAR1−/− (CD45.2+) or WT (CD45.1+) bone marrow. Chimeras reconstituted with only IFNAR1−/− bone marrow served as negative controls for IFNAR1 expression and alloimmunization. (A–B) Representative reconstitution analysis of peripheral blood lymphocytes from (A) muMT− x WT and (B) muMT− x IFNAR1−/− chimeras. Numbers on plots indicate percent of live (left), CD45.2+ (middle), or CD45.1+ (right) cells within the drawn gate. (C) IFNAR1 expression on CD19+ B cells from indicated chimeras. (D) Serum anti-KEL IgG of indicated chimeras 28 days following transfusion with KEL RBCs. Representative of 3 independent experiments with 4–5 mice.
per group. **p<0.01 and n.s., not significant, by Mann Whitney U test. Data from repeated alloimmunization experiments are shown in Supplemental Figure 2C.
Figure 5. IFNα/β promotes germinal center B cell development

IFNAR1−/− and WT mice were transfused with KEL RBCs. (A, C, E) Representative flow cytometric analysis of spleen GC B cells (CD19+IgDloFas+GL7hi) from (A) naïve or transfused mice (C) 8 or (E) 36 days after transfusion. Plots are gated on live CD19+IgDlo cells. Numbers on plots indicate the percent of CD19+B cells within the drawn region. (B, D, F) GC B cells quantified as percent of CD19+B cells (left) and cell number (right). (G) Serum anti-KEL IgG measured by flow cytometric cross-match 36 days after transfusion. (E–G) Mice received a second transfusion 28 days following the first transfusion. Data are from one of two independent experiments with 3–5 mice per group. *p<0.05, **p<0.01, n.s., not significant, by Mann Whitney U test. Data from a repeated independent experiment are shown in Supplemental Figure 3.
Figure 6. IFNα/β promotes plasma cell differentiation

(A–C) WT and IFNAR1<sup>−/−</sup> mice were transfused 28 days following an initial transfusion, and plasma cells were analyzed 14 days later. (A) Representative flow cytometric analysis of plasma cells (CD19<sup>+</sup>IgD<sup>lo</sup>B220<sup>lo</sup>CD138<sup>+</sup>) from bone marrow of naïve and transfused mice. Plots are gated on live CD19<sup>+</sup>IgD<sup>lo</sup> cells. Numbers on plots indicate the percent of CD19<sup>+</sup> B cells within the drawn region. (B–C) Bone marrow plasma cells were quantified as (B) percent of CD19<sup>+</sup> B cells and (C) cell number. Data show one of two independent experiments with 4 mice per group. *p<0.05 by Mann Whitney U test. Data from a repeated independent experiment are shown in Supplemental Figure 4.

(D–F) B cells were isolated from spleens of WT and IFNAR1<sup>−/−</sup> mice by magnetic cell separation. Cells were cultured for 72 hrs in the presence of anti-CD40 +/- rIFNα. (D–E) Representative flow cytometric analysis of live (D) CD19<sup>+</sup>IgD<sup>lo</sup> B cells and (E) B220<sup>lo</sup>CD138<sup>+</sup> plasma cells gated on regions drawn in D). Numbers on plots indicate the (D) percent of live cells or (E) the total number of cells in the drawn regions. Representative of 3 independent experiments. (F)
Summary data of the number of plasma cells generated in the indicated cultures. **p<0.01 by unpaired 2-tailed t-test.