Basophils and IgE: Linking the Allergic Environment to Autoimmunity

Nicolas Charles† and Juan Rivera*

Laboratory of Molecular Immunogenetics, National Institute of Arthritis and Musculoskeletal and Skin Diseases, National Institutes of Health, Bethesda, Maryland 20892, USA

Abstract: As outlined in some of the accompanying articles in this issue, the role of the basophil as an effector cell in allergy and in host defense (particularly to parasites) has long been recognized. However, recent advances advocate for the basophil as an immunomodulatory cell that can promote naïve CD4⁺ T cell commitment to Th2 cell differentiation. While this is in keeping with the concept that the basophil is important in an allergic environment, new discoveries suggest that basophils may be immunomodulatory beyond the context of allergic disease. Here we mainly discuss our own work, which provides a new paradigm for the role of basophils beyond allergy. Our findings demonstrate the importance of autoreactive IgE’s, IL-4 and basophils in promoting autoantibody production and the development of lupus nephritis. The conclusions drawn are based on studies in a mouse model (Lyn⁻/⁻ mice) of spontaneous systemic lupus erythematosus (SLE)-like disease as well as from analysis of the relationship between disease activity in SLE patients and their levels of autoreactive IgE’s and activated basophils with antigen presenting capability. The findings demonstrate a link between the Th2 environment and autoimmunity and provide new areas of investigation with therapeutic potential.

Keywords: Allergy, Basophils, IgE, Lupus, Lyn.

BASOPHILS: ARE THEY A LINK BETWEEN THE ALLERGIC ENVIRONMENT AND AUTOIMMUNITY?

Associated cellular and molecular mechanisms described in parasite infections and allergic disorders have demonstrated the fundamental importance of Th2 cell differentiation and function in these diseases [1, 2]. Indeed, Th2 cytokines (like IL-4, IL-5, IL-13, IL-6, IL-10, TSLP, etc.) and Th2 immunoglobulins are associated in mouse (IgA, IgG1 and IgE) and humans (IgA, IgG4 and IgE) with some parasitic infections and allergies [2]. In both mouse and human, the basophil has been linked to allergic disease [3, 4], with the demonstration in mouse models of a key role for these cells in chronic allergic inflammation [4] and in IgE-mediated systemic anaphylaxis [5]. The recognition that mice, like humans, have circulating basophils has ushered in a new era for this long ignored cell type. Mouse models have allowed the exploration of basophil function in vivo as well as the investigation of the molecular factors important in basophil differentiation and function. As outlined in more detail in other chapters of this issue, recent advances (reviewed in [2, 6, 7]) have made it clear that basophils can cause Th2 cell differentiation in vivo thus demonstrating that these cells have an immunomodulatory role in addition to the long recognized effector role in allergic disease.

For a number of years, we investigated the molecular mechanisms that underlie the function of the high affinity IgE receptor (FcεRI), which is expressed on mast cells and basophils [8]. This effort has led us to explore the role of Lyn kinase, the kinase that phosphorylates FcεRI and initiates signal transduction, in mast cell and basophils [8, 9]. Interestingly, our efforts [10-12], as well as those of others [13-17] led to the recognition that a deficiency in Lyn kinase does not necessarily abrogate allergic responses and, in fact, these mice are hypersensitive to an allergic challenge [10, 18-20]. Lyn kinase is a member of the Src family of protein tyrosine kinases (SFK) and while expressed in most hematopoietic cells it is not expressed in T cells [21]. Nonetheless, the hypersensitivity of Lyn-deficient mice to allergic stimuli provided a strong indication that these mice showed an immune system imbalance whereby Th2 responses were highly favored. Studies characterizing the immune cell characteristics and distribution in Lyn-deficient mice demonstrated that these mice had a constitutive induction of naïve T cell differentiation to Th2 cells [22]. This was dependent on the presence of IL-4 and IgE in these mice but was shown not to be dependent on the presence of mast cells. This led us to explore basophils and whether they might play a role in the observed Th2 skewing. Lyn deficient mice were found to develop a marked peripheral basophilia, with as much as a four fold increase in circulating basophils in unchallenged mice. Importantly, we discovered that an in vivo depletion of the basophils led to a complete stop in the differentiation of naïve T cells to Th2 cells [22]. Collectively, our findings argued that basophils and IgE were key components in the Th2 skewing of Lyn-deficient mice and this axis was essential to the exacerbated allergic phenotype seen in these mice.

It is also well known that in late life Lyn-deficient mice develop a disease that resembles human systemic lupus erythematosus (SLE) [18, 19]. The features of this disease include increased plasma cell numbers, the production of autoantibodies, the presence of vast immune complex deposition in the kidney, and the development of an inflammatory lupus nephritis-like disease that ultimately leads to renal failure. This late life phenotype (usually >20 weeks of age)
is preceded by the allergenic-like (Th2) phenotype (which occurs within 4-6 weeks of age). Thus, this presented an opportunity to explore whether the Th2 phenotype can contribute to the development of a lupus-like disease. While Th2 responses have been described in a lupus nephritis [23], the presence and the role of these responses is unexplored and the concept of Th2 contribution to SLE is controversial, given that Th1, Th17, and loss of regulatory T cell function have been involved [24-30].

In this brief review, we describe new findings that argue for a Th2 component in the development of lupus nephritis. We detail our findings in the context of what is known for the role of Th2 responses in human SLE. Most importantly, and consistent with previous studies [31-35], the findings suggest that basophils are key regulators of humoral immune responses and that dysregulation of their function can have consequences that go well beyond the development or exacerbation of allergic disease.

THE IMMUNOMODULATORY ROLE OF BASOPHILS IN TH2 CELL DIFFERENTIATION AND ALLERGIC RESPONSES.

While this topic will be covered in depth in the accompanying articles, here we briefly outline the major points in basophil regulation of Th2 cell differentiation in order to discuss our findings in the context of the previous work [31-35].

Th2 type cytokines, like IL-4, are known to induce the differentiation of naïve CD4+ T cells into Th2 cells that also produce IL-4, through the induction of transcription factors such as STAT6 and GATA-3 [36]. Th2 cells play an important role in host defense to some extracellular pathogens (parasites or bacteria) and in humoral immune response, but at times they can also have deleterious effects such as in the case of allergies or asthma [36]. Th2 cell responses dampen the ability to induce Th1 cell differentiation and in some cases, like in autoimmunity, this may be beneficial [37]. While it is clear that naïve CD4+ T cells can differentiate into Th2 cells [38], the cellular origin of the IL-4 inducing this commitment is still controversial. The IL-4 derived from the Th2 cell is able to amplify Th2 differentiation [38], but the initial IL-4 signals must come from another cell type that is able to distinguish a Th2-inducing challenge from others. Dendritic cells (DCs) are innate immune cells that have been shown to induce naïve CD4+ T cell to commit to Th2 differentiation, in an IL-4 independent manner, both in vitro and in vivo [39]. However, it is also well known that DCs are not very potent IL-4 producers [39]. Thus it appears that their immunomodulatory function may function in collaboration with other cells that are also able to produce IL-4 and thus induce Th2 cell differentiation.

In humans and mice, basophils are known to produce copious amounts of Th2 cytokines, particularly IL-4, IL-6, IL-13, and TSLP [3]. Basophils can produce high amounts of IL-4 to various stimuli, including FcεRI stimulation, IL-3, and IL-33 treatment [3, 40]. This has led to the thinking that basophils might play a role in the initiation or amplification of Th2 cell differentiation. However, the absence of appropriate tools and information on basophil biology and function, and the slow recognition that mice also have circulating basophils, has hampered their study in vivo. Recent reports [34, 41] have described new tools facilitating the study of basophil involvement in health and disease. Identification of cell surface markers and the development of methods to deplete the basophil in vivo through use of specific antibodies allowed a considerable advance. Antibody-mediated basophil depletion can be accomplished by in vivo injection of antibodies recognizing either CD200R3 (clone Ba103) or the high affinity IgE receptor alpha chain (FceRIα, clone MAR-1) [34, 41]. The basophil depletion is thought to occur via cytolysis through complement activation. However, the precise mechanism still needs to be uncovered. Moreover, what secondary effects might occur after these injections are unknown. Nevertheless, the antibody-mediated depletion of basophils does not deplete mast cells (which also express the FcεRIα) and does not induce detectable changes in the serum levels of cytokines [34, 41]. Such methods are highly useful to analyze the role of basophils in short term immunologic reactions.

These methods have proved quite useful in studying the immunomodulatory role of basophils in allergic reactions [34, 41]. Papain immunization of mice is known to induce Th2 cell differentiation and promote papain-specific IgE and IgG1 production, hallmarks of a Th2 response [34]. Using the approach of antibody-mediated depletion of basophils, Sokol et al. demonstrated that basophils were key players in Th2 cell differentiation induced by subcutaneous injection of papain. Both IL-4 and TSLP produced by basophils were found to be key factors in promoting Th2 cell differentiation [34]. IL-4, IL-6 and IL-13 are also involved in a wide variety of immune regulation, including B cell immunoglobulin (Ig) switch recombination. These cytokines are known to facilitate B cell switch to IgA, IgG and IgE [42-44]. Moreover, IL-4 and IL-6 are known to induce B cell maturation towards memory B cells and plasma cells [43, 44]. Denzel et al. recently showed that basophils were significant contributors in memory humoral immune responses by their ability to secrete IL-4 and IL-6 [35]. After sensitization, basophils were able to induce B cell maturation and lead to an efficient memory humoral response upon a second encounter with the antigen. Thus, this study demonstrated another key immunomodulatory role for basophils as contributors to humoral memory responses [35].

Additional work from the laboratory of H. Karasuyama has also shown an immunomodulatory role for basophils in promoting IgE-dependent chronic allergic inflammation (IgE-CAL) in a mouse model [4, 41]. Basophils were shown to be responsible for promoting the chronic phase of allergic inflammation, including the recruitment of other immune cells type [4, 41]. However, in more acute allergic responses, the effector function of basophils seems dominant as demonstrated by their key role in IgE-mediated systemic anaphylaxis through the release of large amounts of platelet activating factor (PAF) [5]. Thus, it seems that the basophils effector or immunomodulatory roles may be dominant depending on the immune challenge. One factor that may be key in distinguishing the effector from the immunomodulatory phase of basophils may be their ability to present antigen. Three recent studies demonstrated that basophils can process antigen and present it via upregulation of class II major histocompatibility complex [31-33]. In the peripheral blood, basophils are not known to express MHC-II, making this a
Basophils are not known to express MHC-II, making this a feature for their recognition in human peripheral blood [45]. Thus, in the periphery where they can play a role in acute IgG-mediated anaphylaxis, expression of MHC-II is not apparent. Yoshimoto et al., Sokol et al. and Perrigoue et al., showed that human and mouse basophils upregulate MHC-II expression in the secondary lymphoid tissues where they can serve as professional antigen presenting cells in the context of Th2 cell responses. These features make the basophil an important player in the immune system as both an effector and a regulatory cell that can initiate Th2 cell differentiation [31-33] and amplify humoral responses [35].

THE BASOPHILS BEYOND ALLERGY

Like mast cells, the basophils role in the immune system has been mainly focused on its role in acute models of allergic reactions or on chronic models of parasitic infection where they appear to be important upon a second infection with the parasite [46]. However, if the basophil functions as an immunomodulatory cell, one might predict that there would be instances where the basophil might play an important role in immune diseases other than allergic disease. We and others [22, 47] recently reported that genetically modified mice, namely IRF2−/− and Lyn−/− mice, develop a peripheral basophilia and have a constitutive Th2 skewing of the CD4+ T cells. Both genes appear to play a negative regulatory role in peripheral basophil proliferation. The mechanism(s) leading to the constitutive Th2 skewing in vivo was associated with basophils, based on their numbers or on their increased sensitivity to different in vivo stimuli (like IL-3 and IgE-dependent stimulation). As IRF2−/− and Lyn−/− mice showed the development of skin inflammation [48] or of an SLE-like autoimmune disease [18, 19, 49], respectively, these models offer the unique opportunity to explore the role of basophils beyond allergic disease.

BASOPHILS AND AUTOIMMUNE DISEASES

The involvement of basophils in autoimmune disease has been documented in autoimmune urticaria where the development of autoantibodies to FcεRIα have been described and shown to induce basophil activation [50-52]. In atopic dermatitis and other allergic-related skin diseases, basophils and mast cells are known to be involved in promoting the allergic environment leading to allergic cutaneous disease. Nonetheless, there is growing awareness that mast cells can contribute to autoimmune arthritis in some mouse models [53]. In this particular disease, as well as in most of the autoimmune diseases, the effector and immunomodulatory function of the mast cell leads to the initiation of inflammation. The enhancement of inflammation in autoimmune disease, which is the most deleterious component causing tissue damage upon its persistence, can be attributed to Th1 and Th17 cell responses. These CD4+ T cell subsets are believed to be the main effector cells involved in chronic inflammation during autoimmune diseases [54]. As always with such blanket categorization, one is tempted to think that other cell types are not involved in these diseases. However, as mentioned for urticaria and atopic dermatitis, IgE and its high affinity receptor FcεRI, which is expressed on mast cell and basophils, are known to be involved in these diseases. In these aforementioned diseases, a Th2 component is well described and the basophil through its immunomodulatory and effector responses contributes to the development of the autoimmune symptoms. While IgE is clearly recognized as a “Th2 marker” in human pathologies there have been multiple reports [23, 55, 56] where autoimmune patients are described to have increased levels of IgE but are studied in the context of their Th1 and Th17 responses [57]. While the importance of Th1- and Th17-driven responses in the development of autoimmunity is evident, it also seems unlikely that the presence of a Th2 response would not be contributory to the disease. Moreover, in highly complex diseases such as SLE (which might be characterized as a collection of different immunological disorders) it seems likely that the disease results from any imbalance of immune homeostasis [24-30].

From an epidemiological perspective, a parallel between allergy and autoimmunity is evident. Indeed, the incidence of both types of diseases are similarly increasing in developed countries [58]. As previously mentioned, allergy and autoimmunity share a number of common features. In both cases there can be manifestations that are associated with the production of autoantibodies and circulating immune complexes (CIC) formation [59]. Some studies have revealed a common structure of some allergens and autoantigens [60], with a developing hypothesis that many environmental allergens have common structural conformations with self-antigens [60]. This hypothesis could explain the development of certain autoimmune disease such as the chronic urticaria. However, in other autoimmune diseases (such as SLE) where increased IgE levels have been described, these were not associated with atopic features or increased allergic disease in this population [23]. Thus, high levels of circulating IgE may simply reflect the presence of a Th2 response in a non-allergic setting. Given the recent advances outlined above on the role of basophils in Th2 cell differentiation, this could suggest the contribution of basophils in autoimmune pathologies.

Recently it was reported that IgD-mediated basophil activation is relevant to some immune disorders such as HIDS (hyper IgD syndrome), TRAPS (TNF-receptor associated periodic syndrome) or MWS (Muckle-Wells syndrome), all of them being chronic autoinflammatory diseases [61]. The findings suggested that IgD-mediated basophil activation (through a receptor yet to be identified), both in humans and mice, was involved in B cell homeostasis and antibody production, particularly IgA and IgG [61]. Thus, in this context, the basophil has the potential to be involved in autoimmune disorders, where autoantibodies and abnormal levels of IgA, IgG and IgE are found. Indeed, beyond the B cell abnormality and a subsequent humoral dysregulation seen in autoimmune disorders like SLE, the presence of Th2 type Ig’s like IgEs strongly argues for the potential role of an IL-4 secreting cell (Th2 type cell) that influences the humoral response in this pathological setting.

THE LYN DEFICIENT MOUSE MODEL: FROM ALLERGY TO AUTOIMMUNITY

As previously mentioned, Lyn kinase is expressed in most of the hematopoietic cells but not in T cells [21]. The activity of this kinase Lyn is required downstream of various
receptors including cytokine receptors (such as IL-3 and IL-4), tyrosine kinase receptors (such as CD117 or c-Kit), B cell antigen receptor and Fc receptors [62]. As the kinase responsible for phosphorylation of FcεRI, one might have predicted that ablation of its expression or activity would prevent mast cell activation. However, we and others demonstrated that Lyn kinase has dual function as positive and negative regulator of FcεRI signaling [10, 62]. Thus, in the context of increased expression of Lyn kinase (such as seen in mast cells from 129Sv or Balb/c mice), the absence of Lyn causes increased mast cell responses and exacerbation of allergic disease [12, 63]. Interestingly, Lyn kinase also seems to control mast cell proliferation or survival in vivo, since Lyn-/- mice have an increased number of peritoneal mast cell, which is IgE-dependent and is seen early in life [10, 64]. Within 5-6 weeks of age Lyn-/- mice showed increased responsiveness to bronchial challenges (in an asthma model) and to passive systemic anaphylaxis [10, 20]. Lyn-/- mice also have elevated levels of IgM, IgA and IgE in their serum, and this phenotype also develops in early life and is associated with a profoundly altered peripheral B cell compartment and increased numbers of plasma cells [18, 19]. The B cells from these mice were described to be hypersensitive to IL-4 and CD40L stimulation [65], suggesting a mechanism for the strong Th2 responses seen in these mice.

The Th2 Skewing of Lyn-/- Mice is IL-4-, IgE-, and Basophil-Dependent

To directly assess if Lyn-/- mice were Th2 skewed we analyzed CD4+ splenocytes (in the absence of any stimulation) from unchallenged mice and found a large number of IL-4 producing CD4+ T cells, confirming a strong constitutive Th2 skewing [22]. Given the allergic bias of these mice, we set out to explore the role of IL-4, IgE and mast cells in the constitutive Th2 bias by generating mice deficient for both IL-4 and Lyn (Igh7-/-Lyn-/-), IgE and Lyn (Igh7-/-Lyn-/-), and mast cells and Lyn (WSh/WShLyn-/-). Strikingly, mice that were deficient in Lyn and IL-4 (Igh7-/-Lyn-/- (IL-4-/-Lyn-/-)), IgE and Lyn (Igh7-/-Lyn-/-), and mast cells and Lyn (WSh/WShLyn-/-) showed no Th2 skewing. In contrast, mice that were deficient in Lyn and mast cells (WSh/WShLyn-/-) that had a similar constitutive Th2 skewing as in Lyn-/- mice. This suggested that the constitutive Th2 bias required both IgE and IL-4 but not mast cells. This led us to postulate a possible role for basophils, given that they respond to an IgE/Ag stimulus and produce large amounts of IL-4 [2, 31]. This assumption was also in light of the previously mentioned observation that Lyn-/- mice had a peripheral basophilia, with increased number of basophils present in the blood, spleen and peritoneum [22]. Ex vivo FcεRI stimulation of Lyn-/- and WT blood basophils demonstrated a marked increase in IL-4 production on a per cell basis by Lyn-/- basophils [22]. To test Lyn-/- basophil function in vivo, we took advantage of the previous observation that the basophil is a key initial responder upon papain immunization [34]. Strikingly, in Lyn-/- mice immunization with papain led to a rapid increase in specific IgE production, which occurred within 7 days of the first immunization and differed from WT mice where specific IgE is normally observed after a second antigen challenge [22]. Interestingly, the control immunization with human serum albumin (HSA) also led to significant production of HSA-specific IgE whereas this antigen does not induce an effective IgE response in WT mice [34]. This demonstrated that the Lyn-/- mice mounted an early and inappropriate Th2 response. To determine if this Th2 bias altered Th1 responses in these mice, we assessed their ability to respond to a Th1 challenge. WT and Lyn-/- mice were infected with *Toxoplasma gondii*, an intracellular pathogen that initiates a host Th1 response and whose clearance is strictly dependent on IFNγ [66]. A week after infection, the levels of IFNγ in the sera of Lyn-/- mice were about one-half of those in WT mice, and by 1 month of infection Lyn-/- mice had a markedly increased accumulation of cysts in the brain relative to WT mice [22].

While the aforementioned findings provided some evidence for a role of basophils in the Th2 bias of Lyn-/- mice, it was all circumstantial. In order to confirm the role of basophils in the constitutive Th2 skewing of these mice, we adopted two distinct approaches. First, basophils were stimulated in vivo via FcεRI in WSh/WShLyn-/- and WSh/WSh mice. We reasoned that since these mice are mast cell deficient and have high IgE levels, the basophils would be saturated by IgE and injection of rat anti-mouse IgE would selectively target the basophil. Also anaphylaxis would not be an issue in mast cell-deficient mice, since IgE-mediated passive systemic anaphylaxis is mast cell dependent [67]. This challenge resulted in a strong Th2 skewing of WSh/WShLyn-/- mice similar to the constitutive Th2 skewing of Lyn-/- mice. For WSh/WShLyn-/- mice, the Th2 skewing was even more exacerbated than in unstimulated mice, with up to 35% of CD4+ splenocytes producing IL-4 (in the absence of re-stimulation with PMA-ionomycin) [22]. These findings demonstrated that FcεRI-dependent stimulation of basophils in vivo resulted in a potent induction of naïve CD4+ T cell differentiation to Th2 cells. Importantly, we did not observe a Th1 response after the in vivo stimulation of basophils, showing that the activated basophil induced a selective Th2 differentiation [22]. To further confirm these findings we adopted the opposite approach of depleting basophils in WT and Lyn-/- mice. Depletion of basophils in vivo from Lyn-/- mice using the basophil depleting antibody MAR-1, as mentioned above, led to the complete reversal of the Th2 skewing seen in these mice. Mice treated in this manner showed normal CD4+ naïve T cells and no presence of Th2 cells in the spleen. The constitutive Th2 skewing was also reversed in WSh/WShLyn-/- mice by depleting basophils from these mice. Thus, the results described in this section demonstrate that the peripheral basophilia seen in Lyn-/- mice leads to the constitutive Th2 cell differentiation seen in these mice and the presence of IL-4 and IgE is required for this effect [22].

Linking the Th2 Environment to Autoimmunity

It is well known that SLE is a multifactorial autoimmune disease affecting multiple organs. This autoimmune disease affects mainly women in childbearing age (15-40 years old) and can be lethal when kidney involvement (lupus nephritis) is severe and non-responsive to treatment [68]. The etiology of SLE is unknown but it is clear that both environmental and genetic factors play a role in the development of the disease [68]. The American Rheumatology Association (ARA) established 11 distinct criteria to guide physicians in the diagnosis of SLE, based on the clinical manifestations in patients [69]. Presence of at least 4 of the 11 criteria is needed for a definitive diagnosis. Among the criteria is the produc-
tion of autoantibodies to double stranded DNA (dsDNA) and anti-nuclear antigens (ANA) [69], a feature found in other autoimmune diseases. Like in the mouse, these autoantibodies cause the formation of circulating immune complexes (CIC), deposits in the kidney and the chronic inflammation that leads to tissue damage and renal failure. Treatment of SLE currently relies on non-specific immunosuppression and steroids, making new therapeutic approaches of particular interest.

Recently, a polymorphism in an untranslated region of the gene encoding Lyn kinase has been associated with some population of SLE patients and was specifically shown to be correlated with increased autoantibody production [70]. Expression of Lyn kinase was also shown to be lower in the peripheral B cell compartment of some SLE patients relative to normals [71]. Thus, Lyn kinase seems to play a role in SLE and it is not surprising that in late life (after 20 weeks of age) Lyn⁻/⁻ mice develop an autoimmune phenotype that shows many of the features of human SLE. Lyn⁻/⁻ mice develop high levels of autoantibodies specific for double stranded DNA (dsDNA) and nuclear antigens (ANA) with resulting high amounts of circulating immune complexes (CIC) in their serum [18, 19]. Like in human SLE (see below), CICs of IgG- IgA- and IgM- subclasses are deposited in the glomeruli of the kidney of Lyn⁻/⁻ mice leading to the development of a chronic lupus nephritis-like disease, increasing tissue injury, loss of kidney function, and death as mice age [18, 19]. This suggested that the Lyn⁻/⁻ mice could be a suitable model for exploring the effect of a Th2 environment in the development of a lupus-like disease. In particular, given the early life Th2 skewing in these mice and the late life development of SLE in Lyn⁻/⁻ mice independent of genetic background, the argument could be made that the constitutive Th2 environment was permissive to a disease thought to be primarily mediated by Th1 and Th17 cells.

Development of Th2-driven Lupus Nephritis

Aged Lyn⁻/⁻ mice (over 30 weeks old) showed a constitutive Th2 skewing without any apparent increase in Th1 responses as already described in younger mice [22]. At this late age, Lyn⁻/⁻ mice have hallmarks of an autoimmune lupus-like disease with remarkably high levels of autoantibodies and kidney disease [18, 19, 49]. The penetrance of disease was complete with all Lyn⁻/⁻ mice developing some feature of autoimmunity. To investigate if the Th2 environment contributed to the late life autoimmune phenotype of the Lyn⁻/⁻ mice, we studied the SLI-like phenotype of the Ig²⁻/⁻Lyn⁻/⁻ and Il-4⁻/⁻Lyn⁻/⁻ mice. These mice developed nearly all of the features of Lyn⁻/⁻ mice, showing an altered peripheral B cell compartment, splenomegaly, hyper serum IgM and IgA concentrations, and importantly peripheral basophilia. The only difference observed in the Ig²⁻/⁻Lyn⁻/⁻ and Il-4⁻/⁻Lyn⁻/⁻ mice relative to Lyn⁻/⁻ mice was the absence of a constitutively activated Th2 skewed CD4⁺ T cell population [22]. Scoring of the glomerulonephritis (based on cellularity, mesangial proliferation, crescent formation and necrosis) in the Ig²⁻/⁻Lyn⁻/⁻ and Il-4⁻/⁻Lyn⁻/⁻ mice relative to Lyn⁻/⁻ mice revealed that the former mice showed no obvious kidney disease. Evaluation of IgM-, IgA- and IgG-containing immune complexes in their glomeruli showed a marked reduction in their deposition in the kidney and this was associated with relatively little proteinuria. Thus, it was evident that the development of the lupus-like disease in Lyn⁻/⁻ mice was dependent on the presence of IL-4 and IgE; whose absence reversed the Th2 skewing seen in these mice. Strikingly, the titers of IgG anti-dsDNA and ANA as well as the amounts of CIC were also decreased in Ig²⁻/⁻Lyn⁻/⁻ and Il-4⁻/⁻Lyn⁻/⁻ mice, showing that autoantibody production in Lyn⁻/⁻ mice was, at least in part, dependent on the Th2 environment.

These findings suggested the involvement of basophils in autoantibody production, since we previously showed their role in promoting the Th2 skewing of Lyn⁻/⁻ mice [22]. To confirm the involvement of basophils in the production of autoantibodies, we again used the antibody-mediated depletion of basophils to test their role in the production of autoantibodies. Since the depletion of basophils by this approach is transient (6-8 days), and the antibody half-life is normally 2 weeks, the finding that dsDNA and ANA IgG titers were markedly reduced (in some cases by greater than 50%) in 40 week old Lyn⁻/⁻ mice was unexpected. Even more striking was the finding that in younger (20 week old) WSh/Wsh⁻/⁻Lyn⁻/⁻ mice, basophil depletion resulted in an almost complete loss of autoantibody production to levels seen in the control WSh/Wsh mice. One might argue that the observed decrease of autoantibodies might simply reflect the rapid clearance of immune complexes which have a much shorter serum half-life than an uncomplexed antibody [72]. On the other hand, with the continuous production of autoantibodies in autoimmune diseases, one might argue that the marked reduction in autoantibody titers could not have been solely due to the clearance of immune complexes. Analysis of plasma cell numbers in the spleen revealed that the depletion of basophils led to a large decrease (over 60%) of plasma cells in the spleen 6 days post basophil depletion, whereas the latter had no effect on plasma cell number in WT mice. Thus, the rapid decrease in plasma cell numbers, upon basophil depletion, is likely to rapidly affect autoantibody titers. This might suggest a role of the basophil-derived IL-6, which is known to induce B cell maturation, proliferation and survival [44], in the autoantibody production occurring in Lyn⁻/⁻ mice. In fact, we found that after FcεRI stimulation Lyn⁻/⁻ basophils were able to produce large amounts of IL-6 relative to basophils from WT mice [22]. It should be noted, that no depletion of bone marrow plasma cells was observed when basophils were depleted. This argues against a direct effect of the basophil-depleting antibody on the plasma cells, but also shows the limitation of short term depletion on long term autoantibody production. Collectively, the findings show the contribution of the basophil, IgE, and IL-4 to the maintenance of autoantibody production in Lyn⁻/⁻ mice and demonstrate that the Th2 environment can be contributory in autoimmunity.

Activation and Homing of Basophils in Lyn⁻/⁻ Mice

From the above results, it was clear that basophils, IgE and IL-4 were contributory factors in the autoantibody production and the lupus-like disease seen in Lyn⁻/⁻ mice. However, an important question still remained unanswered: How are the hyperresponsive basophils in Lyn⁻/⁻ mice constitutively activated? Because disease development in Lyn⁻/⁻ mice required IgE and IL-4, we postulated that the constitutive activation of basophils might result from the basophil-
independent production of autoreactive antibodies that could cause basophil activation. Interestingly, analysis of autoantibody isotypes revealed high levels of anti-dsDNA and ANA of the IgE class in the sera from Lyn\(^{-/-}\) and W\(^{sh}\)/W\(^{sh}\)/Lyn\(^{-/-}\) mice. This autoreactive IgE was absent or dramatically reduced in the sera of Igh7\(^{-/-}\)/Lyn\(^{-/-}\) and Il-4\(^{-/-}\)/Lyn\(^{-/-}\) mice, respectively. Moreover, IgE-containing CICs were present in the circulation of Lyn\(^{-/-}\) mice. Given that basophils express FcRI, it seemed reasonable to think that these IgE-CICs could be the key factor in promoting basophil activation. Nonetheless, basophils also express Fcy receptors and IgG-CICs were also present in high amounts in the sera of Lyn\(^{-/-}\) and W\(^{sh}\)/W\(^{sh}\)/Lyn\(^{-/-}\) mice. Thus, we explored the ability of IgE or IgG-containing immune complexes (ICs) to induce basophil activation. IgE-ICs were highly effective in activating basophils to produce IL-4 and IL-6 (but not IFN-\(\gamma\), IFN-\(\alpha\), IFN-\(\beta\), TGF-\(\beta\) or IL-12p40), Lyn-deficiency caused secretion of significantly higher amounts of these cytokines relative to WT basophils. In contrast, IgG-containing ICs failed to induce detectable activation of basophils and no cytokine production was observed. These findings argue that the constitutive activation of basophils in Lyn\(^{-/-}\) mice is most likely due to the presence of circulating autoreactive IgE-containing ICs, which are likely to be generated due to B cell hyperresponsiveness to normal levels of IL-4 and CD40 stimulation.

While there are several cell surface activation markers described for human basophils, such as CD203c, CD63 or CD205, these markers are not well defined. Nonetheless, it was important to explore the activation status of basophils in Lyn\(^{-/-}\) mice. CD62L (L-selectin) is a cell surface marker that is expressed on immune cells for their homing to peripheral lymphoid organs and tissues [74] and has been shown to be expressed on human basophils [75]. Measurement of CD62L expression on blood basophils from Lyn\(^{-/-}\) mice showed a marked upregulation of its expression relative to basophils from WT mice. Because expression of CD62L causes homing to lymphoid tissues, we asked if basophils from Lyn\(^{-/-}\) mice could be recruited to lymphoid organs, where they could presumably interact with T and B cells. This possibility might explain how Lyn\(^{-/-}\) basophils are able to induce Th2 skewing of naïve CD4\(^+\) T cells as well as participate in autoantibody production and promote increased numbers of plasma cells. In aged WT control mice, basophils could not be found in lymph nodes. In contrast, in aged Lyn\(^{-/-}\) and W\(^{sh}\)/W\(^{sh}\)/Lyn\(^{-/-}\) mice, basophils were recruited and accumulated in lymph nodes (axial and cervical). As expected, the diminished ability to activate basophils in aged Igh7\(^{-/-}\)/Lyn\(^{-/-}\) and Il-4\(^{-/-}\)/Lyn\(^{-/-}\) mice, the lymph nodes of these mice had only a small percentage (~10\%) of the basophils seen in Lyn\(^{-/-}\) and W\(^{sh}\)/W\(^{sh}\)/Lyn\(^{-/-}\) mice. These findings show that basophils from Lyn\(^{-/-}\) and W\(^{sh}\)/W\(^{sh}\)/Lyn\(^{-/-}\) mice migrate to lymphoid tissues where they can influence both T and B cells. While the production of cytokines, like IL-4 and IL-6, are important towards influencing T and B cells, we also found that basophils from Lyn\(^{-/-}\) and W\(^{sh}\)/W\(^{sh}\)/Lyn\(^{-/-}\) mice express high levels of MHC class II molecules, and high levels of membrane bound BAFF (a B cell survival and activation factor), which has been shown to be associated with SLE [76]. Expression of membrane BAFF did not appear to be dependent on the BAFF receptor, which was detected at very low levels and was not upregulated. Thus, the activation of basophils by circulating IgE-ICs in Lyn\(^{-/-}\) and W\(^{sh}\)/W\(^{sh}\)/Lyn\(^{-/-}\) mice results in the homing of these cells to secondary lymphoid tissues and also causes the upregulation of key surface regulatory molecules that can directly promote T cell differentiation and B cell differentiation and survival. This process is markedly dampened in the absence of IgE or IL-4, as evidenced in Igh7\(^{-/-}\)/Lyn\(^{-/-}\) and Il-4\(^{-/-}\)/Lyn\(^{-/-}\) mice.

The data generated from our mouse studies has allowed us to propose a model (Fig. 1) of how basophils may act to amplify autoreactive responses. We show that in the spontaneous lupus model of Lyn\(^{-/-}\) mice, autoantibodies (dsDNA and ANA) are produced by autoreactive B cells, likely through help provided by autoreactive T cells. How this is initiated is not known, however, the production of autoreactive IgE, likely in response to IL-4 and/or CD40 stimulation, causes basophil activation. IL-4 production by basophils amplifies the production of autoantibodies by promoting increased Th2 cell differentiation and increasing B cell activation and survival, including the survival of plasma cells in the spleen. This is mediated through increased CD62L expression on basophils, which causes their migration and accumulation in peripheral lymphoid tissues, and the cell surface expression of mBAFF and MHC-II expression on basophils in these tissues. Thus, the data from the Lyn\(^{-/-}\) mouse model argues that the basophil is an important contributor in the development and maintenance of lupus-like disease in these mice. Whether this model (Fig. 1) is applicable to the disease seen in other spontaneous models of SLE (like MRL/lpr, NZBW, or BSBX mice) remains to be determined. While most of these models have been shown to have a Th1 and/or Th17 component, it is notable that these mice have been described to have increased levels of circulating IgE [77], suggesting that a Th2 component may contribute to the pathology.

Basophil Dependent Kidney Inflammation

Damage to kidney function in severe lupus nephritis is thought to be irreversible due to the inflammation that causes severe injury and structural alterations to the kidney [78]. Thus, “long term” basophil depletion (over three weeks, with repeated injections of basophil depleting antibody) in 40 week old Lyn\(^{-/-}\) mice did not lead to improvement of renal function (unpublished data). Nonetheless, we could explore whether basophil depletion might alter the inflammatory environment in the diseased kidney of Lyn\(^{-/-}\) mice. Antibody-mediated depletion of basophils was found to cause a marked decrease in pro-inflammatory cytokines detected in the kidney of aged Lyn\(^{-/-}\) mice. Within six days after depletion, a decrease of IL-1β, IL-4, IL-6, and IFN-\(\gamma\) was found with as much as a 70% reduction seen for some cytokines. This demonstrated that basophils (and likely their ability to control autoantibody production) are contributors to the pro-inflammatory environment found in the lupus kidney. Additionally, we found that in the Igh7\(^{-/-}\)/Lyn\(^{-/-}\) mice (which did not have renal failure and showed reduced CIC deposition in the kidney) the pro-inflammatory cytokine environment in the kidney was greatly reduced. Collectively, the findings demonstrate that basophils contribute to the kidney inflammation that occurs in lupus nephritis. Thus, it is of considerable interest to explore if basophil depletion or inactivation,
Basophils and Autoimmunity

The basis of disease in SLE is an escape of autoreactive B and T cells from negative selection (for reasons still not completely understood). These autoreactive lymphocytes lead to the production of autoreactive antibodies “aggregating” with complement factors and autoantigens in the form of circulating immune complexes (CICs). Autoreactive IgGs, IgMs and IgAs being the majority of them, autoreactive IgE, in some circumstances, could accumulate in the CICs. These IgE containing CICs then activate circulating blood basophils leading to their recruitment to lymphoid organs via the CD62L. Once recruited, basophils, via MHC-II and membrane bound BAFF (mBAFF) interact directly with autoreactive lymphocytes. Via their ability to secrete large amounts of IL-4 (and IL-6), basophils then amplify the autoantibody production, leading to the disease amplification.

at an early stage of disease, could delay or rescue the early development of lupus nephritis.

**BASOPHILS AND THE Th2 ENVIRONMENT IN HUMAN SLE**

While mouse models are useful to explore mechanistic and/or physiological aspects that one cannot explore in human disease, the translation of the findings in mouse models to human disease must be explored [63]. Thus, we set out to determine if SLE patients had circulating autoreactive antibodies (anti-dsDNA IgE or anti-IgE IgGs) that might be able to stimulate basophils. As previously mentioned, these autoreactive antibodies were already described to be present in SLE and urticaria’s [23, 50]. However, it was unclear as to whether all SLE patients generated such antibodies. Analysis of plasma samples from a cohort of 43 patients demonstrated the presence of IgE with specificity to dsDNA in all SLE patients screened. When we categorized the patients in 3 distinct groups based on disease activity or SLEDAI score (SLE disease activity index, inactive SLEDAI=0-1, mild SLEDAI = 2-4 and active SLEDAI >4), the levels of anti-dsDNA IgE were strongly associated with increasing disease activity. IgG anti-IgE antibodies were found only in patients with mild or active disease and their levels were also associated with increased disease activity. When patient anti-dsDNA IgE levels were analyzed on the basis of the absence or presence of active nephritis, a striking 4 fold increase in the amount of anti-dsDNA IgE was seen in patients with active nephritis compared to those without active kidney disease. Thus, these findings suggest the direct involvement of autoreactive IgE in human lupus nephritis. Since previous studies [23, 55] had demonstrated that SLE patients had increased circulating IgE, we explored if total IgE was increased in our cohort of patients. While not all patients had increased levels of IgE (whereas all had anti-dsDNA IgE), in patients where total IgE was increased, an association with disease activity and active lupus nephritis was noted. Collectively, the data from human SLE patients demonstrates that beyond the role of Th1 or Th17 cells in this disease, SLE patients also manifest a Th2 component that contributes to increased disease activity and active nephritis.

These findings led us to explore the activation status of blood basophils in SLE patients and healthy (normal) controls. This was based on the assumption that the presence of anti-dsDNA IgE or anti-IgE IgG might lead to basophil activation in these patients. As in the Lyn−/− mouse model, this could cause permanent activation of circulating basophils, via stimulation of FcεRI, leading to amplification of autoantibody production and lupus nephritis. In human basophils, CD203c is known to be expressed upon activation of these cells [79]. Our analysis of CD203c expression on basophils from SLE patients and healthy controls revealed a significant upregulation of this cell surface molecule on blood basophils from SLE patients relative to controls. In addition, as observed in the Lyn−/− mouse model, blood basophils from SLE patients also had increased levels of CD62L on their surface, and the increased CD62L expression correlated with disease activity. This suggested the possibility that, like in the mouse
model, basophils from SLE patients might be found in the peripheral lymphoid organs such as spleen and lymph nodes. Analysis of biopsies from the spleen and lymph nodes of SLE patients and healthy controls revealed a striking difference in basophil homing, with basophils being present at both sites for SLE patients whereas none were detected in the control tissue. In the patient lymph node, basophils accumulated in the periphery or the B cell rich zone of germinal centers, demonstrating that they localize to a site where they could influence both T and B cells. In contrast, we did not find the presence of basophils in kidney biopsies from SLE patients with active nephritis or healthy controls. This could not be analyzed in the Lyn−/− mouse model due to the lack of a suitable reagent for basophil detection in mice. This suggests that the role of the basophil may be immunomodulatory in this disease rather than as a pro-inflammatory effector. Consistent with the homing of basophils to secondary lymphoid tissues, we observed a significant decrease in the number of circulating basophils in SLE patients, and this decrease was associated with increased disease activity. A decrease in the numbers of circulating basophils was not observed in Lyn−/− mice, however, the mice had a peripheral basophilia (driven by the absence of Lyn [22]) and this was not seen in patients. Thus, in the Lyn−/− mouse, the dysregulation of basophil proliferation likely masks the depletion of basophils from the circulation when basophils homed to secondary lymphoid tissues. Unlike the mice, Lyn expression in the basophils of SLE patients was normal when compared to healthy controls.

An alternate explanation for the loss of basophils from the blood of SLE patients could be the immunosuppressive treatment that these patients undergo. While we found that corticosteroids, like prednisone, showed no association with the loss of blood basophils, combination immunosuppressive treatments were associated with decreased blood basophil numbers. However, the immunosuppressive treatments had no apparent effect on the increased basophil activation in SLE patients. This suggests that immunosuppressive treatments may work, in part, through their suppression of blood basophils. This indicates that one might consider the depletion of basophils from the blood of SLE patients as a potential therapeutic strategy. Since the key may well be to diminish basophil activation, another approach might be to deplete IgE (and thus dsDNA-IgE) in SLE patients. This latter approach could be tested rapidly, since the IgE-depleting antibody omalizumab (Xolair®, Genentech), is currently in use for treatment of asthma and allergic rhinitis in some patients and is seemingly well tolerated [80, 81]. Thus, the finding of a role for basophils in SLE [82] may open new avenues with therapeutic potential in the treatment of this disease.

CONCLUDING REMARKS

As described herein, we are at the beginning stages of unraveling the immunomodulatory and effector functions of basophils in health and disease. However, it can be said that beyond their role in allergy and in host defense to parasites, the basophil can promote Th2 cell differentiation and function to amplify the humoral immune response. Its role in autoimmune diseases has not previously been recognized, but the aforementioned findings [82] make a strong argument for a role in lupus nephritis. This role is not in initiation but instead amplification of disease. Key requirements are the autoreactive IgE-dependent activation of basophils, the production of IL-4 by these cells, and the promotion of Th2 cell differentiation. This does not preclude the demonstrated role for Th1, Th17, and the loss of regulatory T cell function, in the development of autoimmunity. Instead, it argues that the strong humoral component seen in autoimmune diseases, like SLE, requires the basophil in order to reach the levels of autoantibody production that result in kidney disease and sometimes kidney failure. Our findings in Lyn−/− mice, as well as in SLE patients, point to the Th2 environment as contributory in SLE.

The recent advances in our understanding of the role of basophils in health and disease demonstrate that a small population of basophil granulocytes can have an important immunomodulatory role in the immune system. However, one must be cautious with the over enthusiastic view of the basophil as necessary for Th2 cell differentiation. To date, experiments dealing with this issue are limited and the key test of challenging mice with a Th2 antigen in the complete absence of basophils remains to be done. Nonetheless, the findings show that in some settings the basophil is a potent inducer of Th2 cell differentiation and through this ability it links the “allergic” environment to the development of an autoimmune disease, like SLE. It will be of interest to explore if these findings will translate to therapeutic benefit.

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