Deletion of the dominant autoantigen in NZB mice with autoimmune hemolytic anemia: effects on autoantibody and T-helper responses

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The mechanisms underlying apparently spontaneous autoimmune diseases, such as autoimmune hemolytic anemia (AIHA) in New Zealand Black (NZB) mice, are unknown. Here, we determine the contribution of the dominant red blood cell (RBC) autoantigen, the anion exchanger protein Band 3, to the development of NZB autoimmune responses. The approach was to prevent Band 3 expression in NZB mice by disrupting the AE1 gene. AE1+/−NZB mice produced RBC autoantibodies at the same levels as the wild-type strain, but they differed in recognizing antigens that correspond to glycophorins, rather than Band 3. Splenic T-helper (Th) cells from wild-type NZB mice proliferated strongly against multiple Band 3 peptides, particularly the dominant epitope within aa861-874. This helper response was severely attenuated in AE1−/− animals, leaving only weak proliferation to peptide aa861-874. The results demonstrate that the defect in self-tolerance in NZB AIHA is directed to the RBC type, and is not specific for, or dependent on, Band 3. However, the predisposition to RBC autoimmunity may be focused onto Band 3 by weak Th cell cross-reactivity between the helper dominant epitope and an exogenous antigen. The redundancy of the major autoantigen illustrates the requirement for specific therapy to induce dominant forms of tolerance, such as T-cell regulation. (Blood. 2007;110:4511-4517)

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Introduction

The underlying mechanisms that drive damaging responses in autoimmune disease are poorly understood. Valuable insights have been obtained from the study of murine models. For example, genetic analyses have shown that different loci govern susceptibility to autoimmunity and determine which organs are affected,1 while functional studies have identified particular autoantigens and highlighted the role of T-cell subsets in pathogenesis.3 However, the stimuli that initiate and perpetuate apparently spontaneous autoaggression remain the subject of controversy.

Autoimmune hemolytic anemia (AIHA) is a classic example of an antibody-mediated autoimmune disease, and provides an opportunity to study the development of specific pathogenic responses, since the major human3-5 and murine6,7 autoantigens have been identified. Efforts to identify possible mechanisms of autoagression often focus on spontaneous AIHA in the New Zealand Black (NZB) strain of mouse. The mice develop signs of anemia by 5 to 6 months of age due to the production autoantibodies that bind red blood cells (RBCs) and cause erythrophagocytosis.8 Band 3, the RBC anion exchanger protein (AE1), is the dominant autoantigen.6,7,9,10 Eluted autoantibodies exclusively immunoprecipitate Band 3 from murine RBCs,6 and 4 of 5 IgG anti–RBC monoclonal autoantibodies (mAbs) derived from NZB mice react with this antigen.8,9,10 The autoantibody response is helper dependent11,12 and Band 3 is also the major target for autoreactive T-helper (Th) cells.7,13-16 Splenic CD4+ T cells from NZB mice, but not H-2–matched healthy strains, proliferate in vitro in response to Band 3,7,15 and initial epitope mapping studies identified a number of stimulatory peptides, of which aa861-874 was consistently the most potent.14 We have now identified the MHC binding and T-cell receptor (TCR) contact residues within Band 3 aa861-874,15 and confirmed that this epitope is dominant in vivo, since the administration of peptides containing the core sequence modulated autoantibody production and disease in NZB mice.16

There are a number of hypotheses put forward to explain the development of autoimmune diseases such as AIHA.17 One widely held belief is that disease is triggered by changes in the presentation of autoantigens from the target cells in such a way as to overcome self-tolerance. For example, infection or inflammation may activate antigen presenting cells (APCs) via innate pathways,18 leading to more effective autoantigen presentation to T cells, or may skew processing pathways within APCs to generate self-peptides that are less subject to central tolerance.19 A second explanation for autoaggression is that the damaging responses are a manifestation of polyclonal activators such as superantigens or mitogens, or of a generalized failure to control lymphoproliferation.20 Another commonly held view is that pathogenic autoimmunity is driven by antigens in the environment, particularly microbial proteins, which cross-react with the target autoantigen.18,19,21 This contention is supported by the induction of murine models, such as experimental autoimmune encephalomyelitis (EAE),22 by immunization with foreign homologues of self-proteins, but there is little evidence for cross-reactivity or mimicry as a cause of apparently spontaneous autoimmune disease.17,23

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dodecyl sulfate–polyacrylamide gel electrophoresis (SDS-PAGE) analysis of wild-type (E) or AE1-knockout (F) murine RBC membrane preparations stained for protein with Coomassie Blue. Each track was loaded with approximately 20 μg protein. The position of Band 3, with an apparent molecular mass of 100 kDa, is indicated (•).

One approach to testing these hypotheses is to determine the contribution of the dominant autoantigen to the induction of disease. Expression of any particular autoantigen in vivo may be unnecessary if disease is driven by aberrant immune processing and presentation of the target cell type, or by polyclonal activation, but would be required if the response were initiated by specific cross-reactivity with an environmental agent. The aim of the current work was therefore to test whether the development of AIHA in NZB mice is dependent on the expression of Band 3, or whether the spontaneous autoimmune response can switch to other RBC antigens. These questions were addressed by disrupting the AE1 gene to prevent Band 3 expression, and then comparing autoantibody and T-cell responses in wild-type and AE1-knockout NZB mice.

Materials and methods

Mice

NZB and healthy control BALB/c mice were maintained under specific pathogen-free conditions in isolators in the animal facilities at the universities of Bristol and Aberdeen. All work was carried out under United Kingdom Home Office Project License approval.

Peters et al.24 provided 129/SV-C57BL/6J strain mice that lack the RBC anion exchanger Band 3 (129/SV-C57BL/6J AE1−/−) due to targeted disruption of the AE1 gene. Failure to express Band 3 prevents the formation of important structural links with the internal membrane skeleton, and results in the production of fragile RBCs of abnormal morphology.24 However, mice homozygous for the null mutation can survive to adulthood, despite failure to express both RBC Band 3 and the related anion exchanger Band 3 (129/SV-C57BL/6J AE1−/−) from Pharmingen (Oxford, United Kingdom). Cell samples containing 10^6 cells were labeled with antibodies at optimal concentrations in the dark for 30 minutes on ice and washed. A benchtop FACScan analyzer IV (Becton Dickinson, Cowley, United Kingdom) was used with a 488 nm argon ion laser to detect green and red fluorescence, dead cells were excluded from analysis by gating on forward and side light scatter properties, and the data were analyzed with MAC Cell Quest (Becton Dickinson).

RBC membrane preparations

Erythrocyte ghosts were prepared by hypotonic lysis of washed RBCs.25 The protein concentration of each sample was determined by the method of Bradford.25

Antigens and mitogen

Murine RBC antigens were prepared and their purity assessed as described previously.7 Briefly, murine Band 3 was purified by anion exchange chromatography from healthy murine RBC membranes that had been stripped of peripheral proteins and detergent solubilized in 1% C_{12}E_8 (Sigma, Poole, United Kingdom). Residual detergent was removed by cold acetone precipitation of the protein, which was dialyzed extensively against phosphate-buffered saline (PBS), pH 7.4, before addition to cultures at a final concentration of 10 μg/mL. The mitogen Concanavalin A (Con A; Sigma), was used to stimulate positive control cultures at an optimal final concentration of 1 μg/mL.

Pepitides

A panel of 92 15-mer peptides with 5 or 6 amino acid overlaps, corresponding to the complete sequence of murine Band 3,26 was synthesized by a multiple-peptide solid-phase method using fluorenylmethoxycarbonyl-polyamide chemistry (Department of Biochemistry, University of Bristol, Bristol, United Kingdom). The purity of the peptides was checked by mass spectrometry. Peptides were used to stimulate cultures at 20 μg/mL, which has been found to be the optimal concentration in previous epitope mapping studies.14,16

Immunization with Band 3 protein

Groups of 6-week-old NZB mice were immunized at the base of the tail by subcutaneous injection of Band 3 protein (100 μg/mouse) administered as an emulsion in complete Freund adjuvant (Sigma).

Culture and stimulation of T cells

As described elsewhere,7,13,16 spleenic T cells were obtained and cultured together in the presence of antigen, peptides, or mitogen in a humidified atmosphere of 5% CO_2 and 95% air at 37°C. On day 5 of culture, proliferation was estimated from the incorporation of tritiated (H) thymidine (Amersham, Aylesbury, United Kingdom) in triplicate samples using a 1450 Microbeta Liquid Scintillation Counter (LKB Wallac, Turku, Finland). Results are presented as stimulation index (SI), representing the ratio of proliferation in stimulated cultures versus unstimulated control wells. As previously, an SI greater than 3 was interpreted as a significant positive response.7,13,16,27

Flow cytometry

The antibodies used were directly conjugated with either fluorescein isothiocyanate or phycoerythrin (anti-mouse CD4, CD8, and CD71) and were purchased from Pharmingen (Oxford, United Kingdom). Cell samples containing 10^6 cells were labeled with antibodies at optimal concentrations in the dark for 30 minutes on ice and washed. A benchtop FACScan analyzer IV (Becton Dickinson, Cowley, United Kingdom) was used with a 488 nm argon ion laser to detect green and red fluorescence, dead cells were excluded from analysis by gating on forward and side light scatter properties, and the data were analyzed with MAC Cell Quest (Becton Dickinson).

Measurement of IgG subclasses bound to RBCs

The number of IgG molecules of each subclass bound to the surface of RBCs was determined using a quantitative and sensitive cellular enzyme-linked immunosorbent assay (ELISA) based on a direct enzyme-linked antiglobulin test (DELAT) developed in our laboratories and described in detail elsewhere.7,16,28 Briefly, a 2% suspension of washed, glutaraldehyde-fixed RBCs was added to duplicate wells of round-bottomed microtiter plates and incubated with subclass-specific sheep anti-mouse IgG antibody (The Binding Site, Birmingham, United Kingdom). After a further incubation with alkaline-phosphatase–conjugated donkey antiserum antibody (Sigma), the RBCs were allowed to react with phosphatase substrate solution (p-nitrophenyl phosphate; Sigma). The supernatant was transferred into flat-bottomed microtiter plates and the optical density (OD) measured at 405 nm (Titerkit Multispec II; Labsystems, Basingstoke, United Kingdom). The number of molecules of each murine IgG subclass bound to...
the RBCs was calculated by interpolation from a standard curve generated by
taking normal RBCs with a known concentration of each purified IgG subclass7,16,28 (Sigma).

**Elution of autoantibody from RBCs**

Autoantibody was eluted from washed RBCs by ether treatment.3,6 Eluates
were stored at −20°C and, after thawing, any precipitated material was
removed by centrifugation.

**SDS-PAGE**

SDS-PAGE was performed in 10% or 12% polyacrylamide gels as
described previously3,6,29,30 according to the method of Laemmli,31 using
the Mini Protean II (Bio-Rad, Hemel Hempstead, United Kingdom).
Apparent molecular masses were calculated from migration relative to
standard protein markers (SDS-6H; Sigma). Where necessary, gels were
developed with Coomassie Brilliant Blue or periodic acid–Schiff (PAS)
stains, respectively, for protein- or glycoprotein-containing bands.32

**Immunoprecipitation of RBC antigens**

Biotin-labeled murine RBC autoantigens or rat antigens were immunopre-
cipitated.3,6,29,30 Briefly, washed RBCs from healthy mice were biotinylated
and sensitized with antibody from sera or RBC eluates. After incubation
with rabbit antiserum against murine IgG Fc (Nordic, Maidenhead, United
Kingdom), the RBCs were lysed in hypotonic buffer and the membranes
dissolved in PBS containing 2% Triton X100 (Sigma). Solubilized immune
complexes were immobilized on sepharose–protein A beads (Sigma) and
eluted into gel sample buffer for SDS-PAGE. Proteins separated by
SDS-PAGE were transferred to nitrocellulose by Western blotting,33 using a
semi-dry apparatus (Pharmacia-LKB, Uppsala, Sweden) with a continuous
buffer system as described by the manufacturer. Biotin-labeled RBC
proteins bound to nitrocellulose were detected by incubation with peroxidase-
conjugated Extravidin (Sigma) and luminoigraphy.3,6,29,30,34 Blots were
stained, respectively, for protein- or glycoprotein-containing bands.32

**Results**

**Generation of AE1−/− NZB mice**

The first step toward determining whether NZB AIHA is dependent
on the dominant autoantigen, Band 3, was to generate genetically
modified AE1−/− mice that lack this protein. A targeted disruption of the
AE1 gene was introduced into 129/SV-C57BL/6J mice,24 back-crossed onto the NZB strain for at least 20 generations, and
the offspring intercrossed to produce AE1−/− NZB mice. It has
previously been demonstrated that mice homozygous for the null
gene express no detectable Band 3 transcript or protein by Northern
analysis. A related question is whether, in the absence of Band 3, the
defect in self-tolerance in NZB AIHA is dependent on Band 3, for example, because it
cross-reacts with an environmental agent, then there should be no
disease in mice that lack the autoantigen. To test these predictions, we first asked whether anti–RBC antibodies are reduced or absent
in NZB mice when the Band 3 autoantigen is not expressed, and
then determined the specificity of any such antibodies.

The levels of RBC-bound IgG were compared in age-matched, wild-type, and AE1−/− NZB mice using a highly sensitive DELAT.28 It
can be seen from Figure 2 that similar levels of all 4 IgG isotypes were
present on the surface of RBCs from both wild-type and BE−/− NZB mice. Although the lack of Band 3 causes hemolysis through increased
RBC fragility,24 autoantibody production in AE1−/− NZB mice is
associated with more severe anemia than is seen in control nonautoim-
mune AE1−/− mice. Mean hematocrit in AE1−/− NZB mice (n = 6) was
11.0% plus or minus 2.5% standard deviation (SD), compared with
19.1% plus or minus 1.5% SD AE1−/− in animals (n = 5) from the
129SV-C57BL/6J strain.

The detection of high levels of RBC-bound IgG in AE1−/− NZB mice raises the question as to the specificity of this immunoglobulin.
A related question is whether, in the absence of Band 3, the genetically modified animals make anti–Band 3 antibodies, which
would be unable to bind their RBCs and therefore remain free in the
serum. To address these issues, sera and antibodies eluted from the
RBCs of AE1−/− NZB mice were tested for the ability to
immunoprecipitate labeled membrane antigens from healthy mur-
ine RBCs (Figure 3). Figure 3A illustrates the electrophoretic
mobility of the major protein (i) and glycoprotein (ii) components
fractionated from wild-type murine RBC membranes. Figure 3B demonstrates that, as previously reported, positive control eluate from the wild-type NZB RBC immunoprecipitated one major protein species (i) corresponding in SDS-PAGE migration to Band 3. In contrast, no antibodies of this specificity were detected in sera (iii,iv) or RBC eluates (v,vi) from AE1^{-/-} NZB mice. However, IgG in these samples instead precipitated a doublet of RBC proteins with apparent molecular masses of 70 kDa to 85 kDa in SDS-PAGE. A comparison with stained RBC proteins and glycoproteins (Figure 3A) reveals that the precipitated bands correspond in mobility either to the internal membrane skeleton proteins Bands 4.1 and 4.2, or to the highly glycosylated cell-surface glycoporphins.32 The diffuse nature of the precipitated bands is consistent with a high degree of glycosylation32 and, of the candidates, only the glycoporphins would be available on intact RBCs for antibody binding.32 To demonstrate that AE1^{-/-} NZB mice retain the ability to make antibodies specific for Band 3, despite the absence of a spontaneous response, a control group of genetically modified animals was immunized with the purified protein as a foreign antigen. Figure 3B (vii,viii) shows that, in addition to the proteins of 70 kDa to 85 kDa, sera from immunized AE1^{-/-} NZB mice also precipitated an RBC membrane component of 100 kDa, corresponding in SDS-PAGE migration to Band 3.

**Characterization of Th responses to Band 3 in wild-type NZB mice**

We have previously reported that splenic Th cells from NZB mice proliferate in vitro in response to Band 3 protein purified from RBCs,7,13 and to a dominant epitope contained within peptide aa861-874.14-16 It was now necessary to generate a complete helper epitope map of Band 3 in wild-type NZB mice with AIHA, for comparison with responses in gene-disrupted AE1^{-/-} NZB mice that lack the autoantigen. A panel of 92 overlapping peptides corresponding to the complete sequence of Band 3 was screened for the ability to stimulate the in vitro proliferation of splenic T cells from NZB mice aged 20 weeks. Figure 4 summarizes the results from a series of up to 8 repeated experiments, and demonstrates that the T cells are capable of responding to a total of more than 20 peptides as judged by a stimulation index of 3 or

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**Figure 3. The spontaneous anti-RBC antibody response in AE1^{-/-} NZB mice targets antigens other than Band 3.** (A) SDS-PAGE analysis of wild-type murine RBC membrane preparations stained for protein with Coomassie Blue (lane i), or for glycoprotein with PAS (lane ii). Each track was loaded with approximately 20 µg protein. Major protein and glycoporphin bands of the RBC membranes are labeled and the apparent molecular masses are indicated in kDa. (B) Immunoprecipitation of biotin-labeled antigens from healthy murine RBCs. Immunoprecipitates were analyzed by SDS-PAGE, blotted onto nitrocellulose, and developed by luminography. Biotin-labeled RBCs were incubated with samples as follows: (i) positive control autoantibody-containing RBC eluate from wild-type NZB mice; (ii) negative control RBC eluate from healthy mouse; (iii,iv) sera from individual AE1^{-/-} NZB mice; (v,vii) RBC eluates from individual AE1^{-/-} NZB mice; (viii,ix) control sera from individual AE1^{-/-} NZB mice immunized with Band 3. The apparent molecular mass of the specific bands precipitated is indicated in kDa. Lanes i, ii, iii, vi, vii, and viii are from different blots.

**Figure 4. Proliferative responses of T cells from NZB mice to a panel of peptides spanning the sequence of the dominant murine RBC autoantigen Band 3, and against the purified protein.** Each bar represents the stimulation index of the peak response of T cells pooled from the spleens from individual mice aged 20 weeks. Each of the 92 peptides was tested on between 5 and 8 mice. The horizontal line indicates the minimum level of a positive response (SI > 3).
above. There were some differences in the profiles of stimulatory peptides between individual mice, but, as reported previously \(^{14-16}\), peptide 87 (aa861-874) invariably elicited strong responses. Purified Band 3 also potently stimulated proliferation in all experiments. Consistent with earlier studies, \(^{14-16}\) responses to Band 3 and peptides were predominantly mediated by T cells of the CD4\(^+\) helper phenotype, as determined by subset depletion experiments, flow cytometric analysis, and the ability of antibodies against MHC class II molecules to block proliferation (results not shown).

**Characterization of Th responses to Band 3 in gene-disrupted AE1\(^{-/-}\) NZB mice that lack the autoantigen**

If autoaggressive Th cells in NZB mice are primed by polyclonal activators or cross-reactive environmental agents, rather than by Band 3, then there should be residual helper responses to Band 3 epitopes in the antigen in vivo. Gene-disrupted AE1\(^{-/-}\) NZB mice, matched in age with the wild-type NZB mice studied in Figure 4, were killed and their splenic Th cells tested for the ability to respond in vitro to the panel of Band 3 peptides and purified antigen (Figure 5). It can be seen that, in comparison with the results obtained using wild-type NZB mice, splenic Th cells from AE1\(^{-/-}\) animals were far less responsive to Band 3 protein and peptides. Thus, AE1\(^{-/-}\) Th cell proliferation against purified Band 3 and the subdominant peptides such as 12 (aa110-125) and 31 (aa301-317) was ablated or very weak. However, although responses to peptide 87 (aa861-874) were also markedly attenuated, significant reactivity to this dominant sequence was retained in all but one of the gene-disrupted animals.

It could be argued that the reduced proliferation against Band 3 epitopes in AE1\(^{-/-}\) NZB mice is not specific, but due to general immunosuppression. However, the mitogen Con A was equally effective in stimulating splenic T cells from wild-type or gene-disrupted NZB mice, and immunization of AE1\(^{-/-}\) NZB mice with Band 3 restored in vitro responses to Band 3 and multiple peptides, including the dominant sequence aa861-874 (results not shown).

The finding that the dominant Band 3 peptide 87 (aa861-874) stimulates residual Th responses in vitro in AE1\(^{-/-}\) NZB mice, albeit more weakly than in the wild-type strain, raises the possibility that the cells are primed in vivo by another antigen. Interrogation of the SwissProt database reveals considerable potential for such cross-reactivity with, for example, multiple bacterial sequences identified that share all or some of the key MHC anchor and TCR contact residues with the autoreactive Th cell epitope in peptide 87. \(^{15}\)

**Discussion**

This study of NZB mice genetically engineered to lack the dominant RBC autoantigen Band 3 has allowed dissection of the underlying mechanisms of spontaneous AIHA. The main result is that induction of NZB AIHA is not dependent on expression of the major autoantigen. In its absence, no detectable antibodies are made to Band 3, and the corresponding helper response is greatly attenuated, but, instead, high levels of autoantibodies with other RBC specificities are generated. The immune defect is therefore specific for the RBC type, rather than for the dominant autoantigen. Such plasticity of the autoaggressive response has implications for the specific treatment of AIHA and other autoimmune diseases.

The shift in the specificity of the autoantibody response in AE1\(^{-/-}\) NZB mice indicates that the potential for autoaggression is not limited to Band 3, and is consistent with the view that a variety of antigens from the target cell may be processed and/or presented in a more immunogenic fashion to drive autoimmune disease. \(^{2,17,19}\) For example, aberrant handling of RBCs by APCs would be expected to predispose to autoimmune responses against a range of potential autoantigens. \(^{17,19}\) Such a change may arise from activation of APCs, or recruitment of nonprofessional APC types, in response to innate immune signals. \(^{17,19}\) Immunoprecipitation experiments revealed that the major autoantigens bound by autoantibody in AE1\(^{-/-}\) NZB mice correspond to the glycophorins, which are heavily glycosylated proteins expressed on the RBC surface. \(^{22}\) Glycophorins are important autoantigens in many other examples of AIHA, including
disease in some human patients,3,4,35,36 dogs,30 and in experimental AIHA induced in mice by immunization with rat RBCs.29,37 Although studies of mAbs from NZB mice support the identification of Band 3 as the dominant RBC autoantigen, they also reveal the potential for autoantibodies with other, minor specificities to be present in NZB AIHA.9,10 Presumably such species, which are not detectable in the polyclonal response of wild-type NZB mice, expand greatly in the absence of Band 3. Autoantibody production in both wild-type and Band 3–deficient NZB mice was associated with anemia. Although the relative pathogenicity of the 2 antibody responses was not compared in detail, the levels bound to RBCs and the proportions of each IgG subclass were similar, and these factors are important determinants of hemolysis.13,16 The correspondence in the ratios of IgG isotypes also indicates that the Th subtype of the responses was unaffected by the switch in specificity.

The lack of anti–Band 3 antibody production in AE−/− NZB mice demonstrates that the response to this antigen in the wild-type strain is not a manifestation of polyclonal B-cell activation by, for example, superantigens.20 This confirms the results of a previous approach to determining the molecular mechanisms responsible for autoantibody production in NZB mice, based on characterization of anti–RBC mAbs derived from NZB splenocytes.9,10,38,39 The IgV H gene sequences of these mAbs, most of which are Band 3–specific,9,10 show evidence of somatic mutation,38,39 indicating that affinity maturation has occurred and that the autoantibody production is therefore antigen driven. The current work allows a further conclusion to be drawn: that the antigen responsible for driving the B-cell response in wild-type NZB mice is Band 3 itself.

Cross-reactivity or molecular mimicry between environmental and autoantigens has often been proposed to be a mechanism underlying apparently spontaneous autoimmune disease.18,19,21,40 Such cross-reactivity may provide increased levels of particular antigenic sequences, or greater avidity of recognition, to overcome self-tolerance, and, in the case of mimicry by infectious agents, innate stimuli may enhance such immunogenicity.18 However, since AIHA in NZB mice is not dependent on expression of Band 3, it is now clear that the disease can develop without the need to invoke any cross-reactivity between environmental agents and this dominant RBC autoantigen. The possibility remains open that the same cross-reactive stimulus initiates responses to both Band 3 and glycoporphin.

Given that Band 3 is not essential for NZB AIHA, and that the response can switch to other autoantigens, the question arises as to why Band 3 is the dominant target in wild-type NZB mice. One explanation is that dominance simply reflects the fact that there are more copies of Band 3 on the RBC surface than any other protein,32 and that the response in AE−/− NZB mice switches to glycoporphins as the next most abundant32 candidate antigens. However, analysis of Th responses in AE−/− NZB mice reveals a new possibility: that cross-reactivity with environmental antigen(s) primes a weak Th response against the major Band 3 epitope, which, in the absence of the autoantigen, cannot spread to other determinants. Such epitope spreading, the recruitment of Th cells with further specificities for the initiating antigen, is a typical feature of autoimmune diseases,2,17,19 and is likely to be important in providing sufficient help to drive autoantibody production41,42 and pathology. In the scenario suggested here, weak cross-reactivity with a single self-epitope acts to focus a tendency toward RBC autoimmunity onto a particular autoantigen, rather than representing a primary or sufficient cause of disease. Several examples of homology between the dominant RBC auto-epitope on Band 3 and bacterial sequences were identified, and it remains to be confirmed whether these, or any other environmental agents, are responsible for priming autoreactive Th cells in NZB mice.

Previous examples of murine genetic modification or mutations that prevent autoantigen expression have been described in other models of autoimmune disease.25 EAE can be induced by immunization with any one of several myelin proteins, and so lack of a particular target, such as myelin basic protein in mutant shiver mice, does not prevent disease induced by the other antigens.25 However, all such models are experimentally induced, and the current study of NZB AIHA is the first to demonstrate redundancy of a major autoantigen in a spontaneous autoimmune disease.

The ability of autoaggressive responses in NZB mice to switch to targets other than the dominant autoantigen is relevant to the development of specific immunotherapy for autoimmune conditions. Thus, even where a single dominant target for pathogenic responses can be clearly identified, reinstatement of tolerance to that autoantigen alone may not be an effective treatment. It will be essential to focus on the induction forms of active tolerance based on regulation,17,42 which may have bystander effects on other associated autoantigens, rather than simply the elimination or silencing of lymphocytes with one set of specificities.

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Authorship

Contribution: A.M.H., F.J.W., and C.-R.S. performed research, analyzed data, and wrote the paper; C.R., L.B., and A.D. collected and analyzed data; S.J.U. and C.J.E. designed research, and wrote the paper; C.R., L.B., and A.D. collected and analyzed data, and wrote the paper.

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