Prevention and partial reversion of the lupus phenotype in ABIN1[D485N] mice by an IRAK4 inhibitor

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ABSTRACT

Objective We have reported previously that the IRAK4 inhibitor PF06426779 given to ubiquitin-binding-defective ABIN1[D485N] mice at 6 weeks of age prevents the major facets of lupus that develop 10 weeks later. The present study was undertaken to investigate whether PF06426779 could reverse the lupus phenotype when administered to 13-week-old ABIN1[D485N] mice that had already developed symptoms of lupus.

Methods Splenomegaly, the number of splenic neutrophils, T<sub>ρ</sub> and Germinal Centre B (GCB) cells, serum levels of immunoglobulins, the extent of kidney, liver and lung pathology, and glomerular IgA and IgM were measured after feeding 13-week-old ABIN1[D485N] and wild-type mice for another 10 weeks with R&amp;M3 diet with and without PF06426779 (4 g/kg).

Results Following drug treatment, spleen size and weight, splenic neutrophil numbers, and serum IgA and glomerular IgA levels of ABIN1[D485N] mice returned to those seen in wild-type mice. The rise in splenic T<sub>ρ</sub> and GCB numbers, the increase in kidney and liver pathology, and the concentrations of serum IgG1, IgG2A and IgE between 13 and 23 weeks were suppressed. There was no reduction in the level of anti-self double-stranded DNA, anti-self nuclear antigens or IgM during the drug treatment.

Conclusions The results demonstrate the therapeutic potential of IRAK4 inhibitors for the treatment of lupus and raise the possibility of monitoring efficacy by measuring decreases in the serum levels of IgA. Our results support the view that there may be a closer connection between lupus and IgA nephropathy than realised previously.

INTRODUCTION

ABIN1 (A20-binding inhibitor of NF-κB 1) is a polyubiquitin-binding protein that restricts the production of inflammatory mediators by the innate immune system. Polymorphisms in TNIP1, the gene encoding ABIN1, predispose to lupus in many human populations.

Key messages

What is already known about this subject?

- Polymorphisms in TNIP1, the gene encoding encoding ABIN1 (A20-binding inhibitor of NF-κB 1), predispose to lupus in many human populations.
- Ubiquitin-binding-deficient ABIN1[D485N] knock-in mice spontaneously develop a lupus-like disease, which is driven by the TR7-MyD88-IRAK4 pathway.

What does this study add?

- The IRAK4 inhibitor PF06426779 reverses splenomegaly, neutrophil numbers and the rise in serum and glomerular IgA levels in ABIN1[D485N] mice that have already developed symptoms of lupus, and prevents further increases in splenic T<sub>ρ</sub> and Germinal Centre B cell numbers, serum IgG1 and IgE concentrations, and kidney and liver pathology.

How might this impact on clinical practice or future developments?

- The study suggests that IRAK4 inhibitors may have therapeutic potential for the treatment of lupus.
- It also indicates that monitoring the serum levels of IgA, as well as patrolling monocyte numbers in the blood, may be a simple non-invasive way to examine drug efficacy, and may reveal whether IgA nephropathy and SLE are the same disease.

expressing the kinase-inactive IRAK4[D329A] mutant or by oral administration of the small molecule IRAK4 inhibitor PF 06426779 prior to the appearance of the hallmark features of lupus, such as glomerulonephritis, liver and lung inflammation, and increased levels of autoantibodies. A different IRAK4 inhibitor (BMS-986126) was also shown to prevent kidney pathology in the lupus-prone MRL/lpr and NZB/W mouse lines.

Lupus is usually diagnosed when humans have already developed the disease. To evaluate the potential of the IRAK4 inhibitor for the treatment of lupus, it was therefore critical to investigate whether it prevented the further rise, or even reversed, the hallmark features of lupus after they had already
developed. Here, we present the results of such a study in ABIN1[D485N] mice, which has revealed some interesting and unexpected findings.

**METHODS**

ABIN1[D485N] mice aged 13 weeks that had already developed splenomegaly, autoimmunity and organ inflammation, or control wild-type littermates, were fed for 10 weeks on chow containing the IRAK4 inhibitor PF06426779 or control chow. The sources of other reagents and the methods used are described elsewhere. Statistical analysis was carried out using GraphPad Prism 9 software. The distribution was determined using the Shapiro-Wilk normality test. Multiple comparisons of data with normal distribution were performed using one-way ANOVA followed by Tukey’s post hoc test. Multiple comparison of non-parametric data was done using the Kruskal-Wallis test, followed by the Mann-Whitney U test. Data in percentages were logit-transformed. Figures were drawn using GraphPad Prism and Adobe Illustrator.

**Immunohistochemistry**

Kidney sections were stained with haematoxylin and either anti-IgA (Abcam, ab97231) or anti-IgM (Abcam, ab97226) primary antibodies at 1:1600 dilution. The signal was detected using ImmPRESS[R] HRP Horse Anti-Goat IgG Polymer Detection Kit (Vector Laboratories). Slides for microscopy were scanned using Motic EasyScan Infinity at ×40 magnification and analysed using QuPath software. Forty glomeruli per mouse were gated and those staining positively with 3,3'-diaminobenzidine were quantified.

**RESULTS AND DISCUSSION**

The IRAK4 inhibitor PF06426277 was given to ABIN1[D485N] mice in their food (4 g/kg) from 13 weeks. At this age, splenomegaly, autoimmunity and organ inflammation have already started to develop. Strikingly, 10 weeks after treatment with PF06426779 had commenced, the spleen size had almost returned to that seen in wild-type mice (figure 1A) and neutrophil numbers had also normalised (figure 1B). Splenic T FH and Germinal Centre B (GCB) cell numbers continued to increase over the 10-week period in ABIN1[D485N] mice not given the drug, but this was prevented by the IRAK4 inhibitor (figure 1C,D). Importantly, the further rise in kidney and liver pathology from 13 to 23 weeks was suppressed, but not lung pathology (figure 2A–F).

ABIN1[D485N] mice display elevated levels of patrolling monocytes in their blood at only 4 weeks, which increases progressively up to 16 weeks. PF06426779 administered from 6 weeks prevented the subsequent rise in pMo seen in the absence of PF06426779. Here, we found that PF06426779 administered from 13 weeks reduced the number of pMo after 18 or 23 weeks, but not to the level seen in control WT mice (online supplemental figure S1).

The kidneys of ABIN1[D485N] mice contain deposits of complement factors and immunoglobulins (IgA, IgM, IgG, IgA, IgM, IgG) that are present in higher concentrations in ABIN1[D485N] mice compared to wild-type control mice. Immunohistochemical analysis revealed that these deposits are not well visualised at 13 weeks, but are more prominent at 23 weeks after IRAK4 inhibitor treatment.

Figure 1  Reversal of splenic phenotypes in ABIN1[D485N] knock-in mice by an IRAK4 inhibitor. 13-week-old ABIN1[D485N] mice (n=6) and age-matched wild-type (WT) (n=5) mice were fed for 10 weeks on R&M3 diet containing or not containing PF06426779 (4 g/kg). Mice were culled at the age of 23 weeks. An additional group of 8 WT and 6 ABIN1[D485N] knock-in mice were culled when they were 13 weeks old as a control. (A) Spleen weight and a representative image showing relative spleen size. Bar equals 1 cm. (B) Neutrophil numbers in the spleen. (C) Germinal Centre B cell numbers in the spleen. (D) Follicular T helper cell numbers in the spleen. Each symbol shows the result from a single individual mouse. Statistical significance was calculated using one-way ANOVA and Tukey’s post hoc test (A, B and D) or the Kruskal-Wallis and the Mann-Whitney U tests (C); * denotes p<0.05, ** denotes p<0.01 and *** denotes p<0.001. The methodology used is detailed in Nanda et al.6
IgM and IgG) at 5–6 months of age. We therefore measured the levels of these and other immunoglobulins in the serum of ABIN1[D485N] mice (figure 3). At 13 weeks, IgA, IgM, IgG2a and IgG2b levels were elevated (figure 3A–C), but IgG1 and IgE were not (figure 3E,F). Treatment of ABIN1[D485N] mice for 10 weeks with the IRAK4 inhibitor reversed the rise in serum IgA seen at 13 weeks of age, the levels returning to those measured in WT mice (figure 3A). In contrast, serum levels of IgG1 and IgE continued to increase between 13 and 23 weeks in ABIN1[D485N] mice whether or not the mice were treated with the IRAK4 inhibitor. In contrast, the further rise in serum IgG2a observed between 13 and 23 weeks in ABIN1[D485N] was prevented by the drug (figure 3D).

Serum IgG1 and IgE levels were not elevated in 13-week-old ABIN1[D485N] mice but were greatly elevated after 23 weeks (figure 3E,F). Similar to IgG2a, this rise between 13 and 23 weeks was prevented by administration of the IRAK4 inhibitor (figure 3E,F).

The lack of effect of the drug on serum IgM and IgG2b levels does not appear to be explained by much longer
half lives of these immunoglobulins in the blood. In normal adult mice, all the immunoglobulins studied have half lives of days and not weeks or months. The half lives of IgM and IgG2b are 2 and 4–6 days, respectively, shorter than IgG1 (6–8 days), but longer than IgA and IgE (1 day and 0.5 day, respectively).

Interestingly, the 10-week treatment with the IRAK4 inhibitor did not affect serum levels of ANA significantly (figure 3G) or anti-dsDNA (figure 3H) observed after 23 weeks. The reversal of splenomegaly and splenic neutrophil levels, the prevention of the further rise in splenic T eff and GCB cell numbers, and the improved kidney and liver pathology over this period following treatment with the IRAK4 inhibitor therefore occurred without any significant change in the serum levels of the autoantibodies examined.

Of all the antibodies studied, the effect of the IRAK4 inhibitor on serum IgA levels showed the strongest correlation with the appearance and disappearance of splenomegaly and splenic neutrophil numbers between 13 and 23 weeks of age. Consistent with these findings, glomerular IgA was also reduced by feeding 13-week-old ABIN1[D485N] mice for 10 weeks with diet containing 06426779, but IgM was not (online supplemental figure S2). Interestingly, the deposition of IgA in the kidney, termed IgA nephropathy (IgAN), is the most frequent primary glomerulopathy in the world. Although the co-association of SLE and IgAN had been reported in several human patients, the possibility that IgA deposition might be a subtype or even a primary trigger of kidney pathology in human SLE had not been considered until a patient diagnosed with SLE was found to contain glomerular IgA deposits. The ABIN1[D485N] mice also accumulate glomerular IgA deposits. Since IgAN and SLE are both relatively common conditions, it is possible that their co-existence is underdiagnosed.

Four IRAK4 inhibitors have entered clinical trials for the treatment of rheumatoid arthritis, psoriasis and lymphoma with little evidence of toxicity. The present study suggests that IRAK4 inhibitors may also have therapeutic potential for the treatment of some types of lupus and that it may be possible to monitor their efficacy rapidly and non-invasively by measuring the serum levels of IgA and/or other immunoglobulins, such as IgG1 or IgE or by a decrease in patrolling monocyte numbers. The routine monitoring of serum IgA levels in patients with lupus patients may also reveal whether IgAN and SLE are really the same disease.
Brief communication

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