**TLR7 gain-of-function genetic variation causes human lupus**

Although circumstantial evidence supports enhanced Toll-like receptor 7 (TLR7) signalling as a mechanism of human systemic autoimmune disease, evidence of lupus-causing TLR7 gene variants is lacking. Here we describe human systemic lupus erythematosus caused by a TLR7 gain-of-function variant. TLR7 is a sensor of viral RNA and binds to guanosine. We identified a de novo, previously undescribed missense TLR7Y264H variant in a child with severe lupus and additional variants in other patients with lupus. The TLR7Y264H variant selectively increased sensing of guanosine and 2',3'-cGMP, and was sufficient to cause lupus when introduced into mice. We show that enhanced TLR7 signalling drives aberrant survival of B cell receptor (BCR)-activated B cells, and in a cell-intrinsic manner, accumulation of CD11c+ age-associated B cells and germinal centre B cells. Follicular and extrafollicular helper T cells were also increased but these phenotypes were cell-extrinsic. Deficiency of MyD88 (an adaptor protein downstream of TLR7) rescued autoimmunity, aberrant B cell survival, and all cellular and serological phenotypes. Despite prominent spontaneous germinal-centre formation in Tlr7Y264H mice, autoimmunity was not ameliorated by germinal-centre deficiency, suggesting an extrafollicular origin of pathogenic B cells. We establish the importance of TLR7 and guanosine-containing self-ligands for human lupus pathogenesis, which paves the way for therapeutic TLR7 or MyD88 inhibition.

Although systemic lupus erythematosus (SLE) is generally a polygenic autoimmune disease, the discovery of monogenic lupus cases and rare pathogenic variants has provided important insights into disease mechanisms, including important roles of complement, type I interferons and B cell survival. There is accumulating evidence that patients with SLE display phenotypes that are consistent with increased TLR7 signalling associated with elevated IgD−CD27− double-negative B cells and, more specifically, the CXCR5−CD11c+ subset (also known as DN2 B cells or age-associated B cells (ABCs)) in the peripheral blood, and excessive accumulation of extrafollicular helper T cells. Genome-wide association studies have identified common polymorphisms in or near TLR7 that segregate with SLE. There is also conflicting evidence as to how TLR7 overexpression causes autoimmunity, particularly, the relative roles of TLR7-driven spontaneous germinal centres (GCs) versus the role of TLR7-driven double-negative B cells; the latter have been proposed to originate extrafollicularly and be pathogenic in lupus. Most mouse lupus models in which TLR7 has a role in pathogenicity display increased formation of GCs and T follicular helper (Tfh) cells, and it has been proposed that TLR7 drives GCs enriched in self-reactive B cells. However, recent reports have demonstrated that lupus can develop independently of GCs in mouse models in which disease is dependent on MyD88 signalling. TLR7 and TLR8 selectively detect a subset of RNA sequences. On the basis of recent knowledge of how TLR8 senses RNA degradation products to trigger downstream signalling, it is thought that one ligand-recognition site in TLR7 binds to guanosine or 2',3'-cyclic phosphoguanosine monophosphate (cGMP), derived from GTP degradation, which synergizes with uridine-rich short RNAs binding to a second site. Here we describe the action of a de novo TLR7...
single-residue gain-of-function (GOF) variant that increases the affinity of TLR7 for guanosine and cGMP, causing enhanced TLR7 activation and childhood-onset SLE.

**TLR7 variants in patients with SLE**

We undertook whole-genome sequencing of a Spanish girl who was diagnosed with SLE at the age of 7 (Supplementary Table 1). She first presented with refractory autoimmune thrombocytopenia and had elevated anti-nuclear antibodies (ANAs) and hypocomplementaemia. She went on to develop inflammatory arthralgias, constitutional symptoms, intermittent episodes of hemichorea, and had mild mitral insufficiency and renal involvement after admission with a hypertensive crisis. Bioinformatics analysis revealed a de novo, TLR7 p.Tyr264His (Y264H) missense variant that was predicted to be damaging by SIFT and CADD (Fig. 1a–c (family A) and Supplementary Table 2). This variant was not present in the databases of normal human genome variation (gnomAD, ExAC, dbSNP). Examination of the BAM files together with paternity analysis confirmed that the mutation occurred de novo (Extended Data Fig. 1a, b, d). The mutated tyrosine residue lies in the eighth leucine-rich repeat of TLR7, within the endosomal part of the receptor (Fig. 1b) and is highly conserved across species, including zebrafish (Fig. 1d). Additional analyses for rare variants in 22 genes that can cause human SLE when mutated (Supplementary Table 3) revealed a heterozygous variant in RNA5E.H28, p.Ala177Thr, which, when homozygous, causes SLE.

Whole-exome sequencing (WES) analysis of additional patients with SLE identified two other variants in TLR7 (B.I.2 F507L and C.I.1 R28G; Fig. 1a–d, Extended Data Fig. 1c, Supplementary Tables 1, 2). Notably, in family B the mother had SLE from her mid-twenties and the daughter was diagnosed with neuromyelitis optica in the presence of ANAs and antibodies to aquaporin-4 (AQP4-Ab, also known as NMO-IgG) in the serum and cerebrospinal fluid. Both carried the TLR7 Y264H variant, which was also highly conserved (Fig. 1d). No additional rare variants in the 22SLE-causing genes were identified in these families (Supplementary Table 3).

**TLR7264H increases guanosine sensing**

To test whether pathogenic variants enhance TLR7 signalling and down-stream NF-kB activation, we transfected RAW264.7 cells with plasmids encoding the different TLR7 mutants. Compared with wild-type cells, overexpression of TLR7264H and TLR7507L constructs showed enhanced NF-kB activation for the mutant variants after 2′,3′-cGMP stimulation, whereas overexpression of TLR7507L showed enhanced NF-kB activation after stimulation with guanosine and single-stranded RNA (ssRNA) (Fig. 1e, f). Using a published TLR7 structure, we mapped the mutated Tyr264 residue to TLR7 ligand-binding site 1, where R837/R848 and Tyr264 residue to TLR7 ligand-binding site 1, where R837/R848 and side-chain OH has been shown to form a hydrogen bond with 2′,3′-cGMP. To further understand whether the Y264H mutation could enhance TLR7 sensing or activation, we used the method of thermodynamic integration within molecular dynamics simulations to determine the relative binding affinity of the ligands to dimeric TLR7 (Extended Data Fig. 1e, f). Guanosine, but not R848, appeared to have increased affinity to its binding site with both singly (H264) and doubly protonated (H+264) protonated variants (Fig. 1g). The site of the mutation, Tyr264, was close (albeit not bound) to the guanosine, which formed a hydrogen bond to Thr286, limiting the access of water to this part of the ligand (Fig. 1h). By contrast, the mutants H264 and H+264, appeared to allow solvent to access this region, helping to stabilize a cluster of water molecules that provide a favourable environment for the polar ribose ring of guanosine (Fig. 1i). In these simulations, Y264H also displayed stronger attractive electrostatic interactions with guanosine, providing a plausible explanation for the much larger predicted affinity to this variant. The electrostatic attraction of H264 is predicted to be even greater for the natural ligand 2′,3′-cGMP that carries a negative charge.

**TLR7264H causes autoimmunity in mice**

Our modelling suggested TLR7264H would increase affinity to endogenous ligands. To investigate whether TLR7264H could cause SLE, we introduced the orthologous allele into C57BL/6 mice using CRISPR-Cas9 editing. WES analysis confirmed that TLR7264H was the only relevant CRISPR-induced coding variant segregating with the phenotype. The resulting strain was named kika and TLR7264H is hereafter named the kik allele. A CRISPR-generated line lacking TLR7 protein owing to a 1-bp deletion was included as a control (Extended Data Fig. 2a, b). Male or female mice (aged 12 weeks) carrying one or two kik alleles displayed splenomegaly with increased cellularity (Fig. 1j), decreased survival (Fig. 1k) and ANAs with nuclear, cytoplasmic, cell-cycle-dependent and Golgi staining (Fig. 1I, Extended Data Fig. 3a). These were detected from 6 weeks of age in female kika mice (Extended Data Fig. 3d, e). Male TLR7507L and female TLR7507L kika mice also developed antibodies to the TLR7 ligands ssRNA and Smith protein (Sm) and ribonucleoprotein (RNP), an RNA-containing nuclear self antigen (Fig. 1m), and 10–20% had weak double-stranded DNA reactivity (Extended Data Fig. 3b, c). These results suggest TLR7264H is an X chromosome-linked dominant GOF allele.

**H264 increases the response to guanosine**

To validate the predicted increased affinity of H264 for guanosine, we tested the response of TLR7264H kika bone-marrow-derived macrophages (BMDMs) to increasing doses of guanosine and R848. Although no difference was observed with R848 (Fig. 1n and data not shown), we found an increased responsiveness to guanosine from kika BMDMs compared with wild-type BMDMs (Fig. 1o and Extended Data Fig. 3g). Both the first (guanosine-binding) and second (uridine-binding) sites of TLR7 are necessary for ssRNA-induced TLR7 signalling. To test the responsiveness of wild-type and mutant BMDMs to ssRNAs lacking uridine (ss41-L) or containing 6 to 10 uridines. Interestingly, whereas ssRNA sensing by wild-type BMDMs correlated with uridine content, ssRNA sensing by kika BMDMs was independent of uridine content, and ssRNA lacking uridine induced TLR7 activity in kika cells (Extended Data Fig. 3h). These results collectively indicate that the kika mutation selectively increases sensing of guanosine to the first site independently of activity at the uridine-selective second site, therefore raising the sensitivity to otherwise non-TLR7-stimulating ssRNAs.

**Tissue damage caused by TLR7264H**

In-depth phenotyping of kika mice revealed marked thrombocytopenia as seen in the proband (Fig. 2a) and a slightly lower white blood cell count (Extended Data Fig. 5a). Proliferative glomerulonephritis was evident in kidneys (Fig. 2b), as well as expanded mesangial matrix with electron-dense deposits and increased mesangial cellularity (Fig. 2c). Lymphoid infiltrates were seen in the liver, salivary glands and pancreas (Fig. 2d) where they occasionally formed peri-islet follicular structures. Exocrine pancreatic tissue was often replaced by fat, particularly in TLR7264H mice (Fig. 2d). Other findings included subpleural, perivascular and interstitial infiltrates in the lungs; myocyte degeneration and necrosis in skin panniculus muscle; focal myocardial fibrosis, splenic lymphomas (4 out of 6 mice); chronic lymphadenitis in the lymph nodes and gut; and hyperplasia of Peyer’s patches (Supplementary Table 4, Extended Data Fig. 4). Serum levels of IFNγ, IL-6, IL-10 and TNF were increased (Fig. 2e, Extended Data Fig. 3f).

Flow cytometry analysis of kika spleens revealed a reduced T:B cell ratio with an increase in total B cells, spontaneous GCs, and increased plasma cells and ABCs; ABCs were also expanded in the blood and kidneys (Fig. 2f–i, Extended Data Fig. 5b, c). The percentages of splenic marginal zone B cells were decreased, but the total numbers were not decreased (Extended Data Fig. 5d). Effector or memory CD4+ CD44high cells, including Tfh cells and CXCR3+ extrafollicular helper CD4+ T cells, were increased in kika mice (Fig. 2j–l) as well as circulating...
CXCR5<sup>hi</sup>PD-1<sup>hi</sup> GC TFH cells that are usually only seen in secondary lymphoid tissues (Extended Data Fig. 5e). Plasmacytoid dendritic cells (Extended Data Fig. 5g), supporting that the expansion of these subsets in kika mice is driven by the TLR7 GOF.

**B-cell-intrinsic effects of TLR7<sup>Y264H</sup>**

To establish which phenotypes were cell autonomous, mixed bone marrow chimeras were generated by adoptively transferring 100% wild-type CD45.2 or kika CD45.2 bone marrow, or 50:50 mixes of either wild-type CD45.1:kika CD45.2 or wild-type CD45.1:wild-type CD45.2 bone marrow into sublethally irradiated Rag<sup>−/−</sup> mice (Fig. 3a, b, Extended Data Fig. 5h). Autoantibodies were present in chimeric mice receiving either 100% or 50% kika bone marrow cells (Fig. 3a, Extended Data Fig. 5j). Expansion of GCs, ABCs and plasma cells (Fig. 3b) was cell-intrinsic whereas all T cell phenotypes and reduction in marginal zone B cells were largely cell-extrinsic (Fig. 3b, Extended Data Fig. 5l, i). The ABCs of kika mice expressed more TLR7 although not to the extent seen in Yaa mice that express two copies of TLR7 (Fig. 3c). By contrast, the functional consequences of the Y264H mutation were more severe than the Yaa allele, as seen by higher levels of anti-RNA and smRNP autoantibodies and ABCs in kika mice (Fig. 3d, Extended Data Fig. 5k).
We investigated whether the Y264H variant leads to spontaneous TLR7 cleavage and activation in the absence of stimulation. Western blot analysis of splenocyte lysates from kika and wild-type littermates using two different antibodies against both the C and N termini revealed the presence of the approximately 75-kDa C-terminal-cleaved and 65-kDa N-terminal cleaved TLR7 product in splenocytes from immunized kika mice (Fig. 3f). Such cleaved fragments have been reported to be indicative of the active form of TLR7. MyD88 was also increased in splenocytes from unimmunized kika mice (Fig. 3e). Such cleaved fragments have been reported for the 65-kDa N-terminal cleaved TLR7 product in splenocytes from unimmunized kika mice (Fig. 3f) and were consistent with enhanced TLR signalling. MyD88 was also increased in splenocytes from unimmunized kika mice (Fig. 3e).

**Enhanced survival of BCR-activated cells**

We next examined the stage at which TLR7 breaks B cell tolerance. We hypothesized that constitutive TLR7 signalling may provide an aberrant signal to self-reactive B cells that have bound to self-antigen through their BCR (signal 1) and would otherwise die within 72 h, as
occurs in anergic B cells and in immature CD93+ cells stimulated with anti-IgM. We could not use CD93 to purify immature splenic B cells because we found that agonistic TLR7 treatment of mature B cells upregulated CD93 (Extended Data Fig. 7a). We therefore activated either total splenic B cells or CD93+B220+ immature bone marrow cells from kika, wild-type and Yaa mice carrying a Tlr7 duplication, stimulated them with R837 or anti-IgM and performed live cell counts 72 h later. We observed that anti-IgM, but not R837, enhanced the survival of total, mature and immature kika B cells compared with control cells (Fig. 3g, Extended Data Fig. 7b–d). This differs from reports using TLR7 transgenic B cells, which displayed increased survival only when activated with a TLR ligand and not with anti-IgM, again suggesting sustained activation of TLR7(Y264H) by endogenous ligands. RNA-sequencing (RNA-seq) analysis of kika and control splenocytes cultured for 20 h with anti-IgM revealed that 203 and 34 transcripts were upregulated or downregulated, respectively, by more than twofold ($P < 0.05$ in kika cells (Fig. 3i). Upregulated transcripts included the anti-apoptotic genes Mxd3 and Serpina3g, as well as Il28ra (also known as Ifnar1), which encodes the common IFN-$\lambda$1/2/3 receptor. Levels of the transcription factor SOXS, which decreases B cell proliferative capacity while allowing plasmablast differentiation, were also high. Other upregulated transcripts included Cxcr3, which promotes lupus nephritis. We confirmed a decreased tendency for apoptosis in kika ABCs with decreased expression of caspase-3 and also a small decrease in proliferation (Fig. 3h). Overall, these results suggest that hypersensitive TLR7 signalling enables the survival of B cells that bind to self-antigen through their surface BCR.
**MyD88 dependence and GC independence.**

To confirm that the observed aberrant B cell survival after IgM stimulation was due to enhanced TLR7 signalling, we crossed kika mice with MyD88-knockout mice. MyD88 deficiency completely rescued kika phenotypes, including splenomegaly (Fig. 4a), accumulation of ABCs, GC B cells, plasma cells, e1* cells (Fig. 4b), Extended Data Fig. 8) and autoantibody formation (Fig. 4c). The aberrant survival of B cells
receiving only signal 1 was completely abrogated in anti-IgM-activated kika B cells lacking MyD88 (Fig. 4d) without changes in proliferation (Fig. 4e).

It remains controversial whether the spontaneous GCs of lupus-prone mice contribute to the autoimmune phenotype, with some suggesting that TLR7 promotes the appearance of self-reactive GC B cells that produce autoantibodies and others proposing that the pathogenic B cells are ABCs of extracellular origin. To resolve this question, we crossed kika mice with B6CBAf2/Cd23mice that cannot form GCs. In the F2, intercross offspring, we enumerated GC B cells and confirmed that kika B6CBAf2/Cd23mice had a substantial reduction in GC B cells (Fig. 4f). Despite the paucity of GC B cells, kika B6CBAf2/Cd23mice developed autoantibodies and an even more pronounced expansion of ABCs and plasma cells than that observed in their GC-forming kika littermates (Fig. 4g, h). These results support the idea that TLR7-driven autoimmunity is GC independent.

We finally analysed the PBMCs from proband A.I.1 carrying the Y264H variant to confirm the phenotypes observed in the kika mouse. Flow cytometry analysis revealed that ABCs and the parental IgD− CD27+ B cells were substantially increased compared with gender- and age-matched controls (Fig. 4j, k). As seen in kika mice, TLR7 and MyD88 protein expression were increased in the pDCs and B cells of the proband (Fig. 4i, m), as was the cleaved TLR7 product (Fig. 4i). Increased levels of ABCs, TLR7 and MyD88 were not seen in the most unrelated patients with SLE (Extended Data Fig. 9a, b). PBMCs in proband A.I.1 also revealed upregulation of the NFκB activation marker CD25 in unstimulated monocytes (Fig. 4n), increasing its expression after TLR7 stimulation (Extended Data Fig. 9c). This variant allele, but not the mother (A.1.2), carrying the RNAMEH2B variant allele (Fig. 4o), had an increased type IFN signature.

We conclude that TLR7 GOF can cause B-cell-driven autoimmunity including SLE due to increased affinity to guanosine, leading to a lowered threshold for TLR7 activation. Although the human TLR7Y264H variant is sufficient to induce lupus in mice with no clear additive effects of Rnaseh2b hemizygosity apart from increased type I IFN gene transcripts, an exacerbating role of this variant in humans may occur in the presence of environmental stimuli, including ssRNA viruses such as SARS-CoV-2 that are dependent on TLR7 immunity. The TLR7 GOF promotes the survival of BCR-activated immature B cells, which are known to be enriched in self-reactivity. Activation of self-reactive B cells by self-antigen in the presence of a constitutive signal 2 is likely to promote differentiation and autoantibody production of B cells that would otherwise be destined to die in the absence of T cell help. Notably, despite enhanced TLR7 signalling causing the cellular accumulation of ABCs and GC B cells, GC B cells are dispensable for the autoimmune phenotype and rather protect against it. Extracellular ABCs are therefore the most likely source of pathogenicity. It will be important to determine whether this is true for all cases of SLE, or only for patients in whom excessive TLR7 signalling is the dominant pathogenic pathway. Although highly damaging TLR7 GOF mutations are rare, our data, together with evidence of increased TLR7 signalling in a large fraction of patients with SLE, suggest that TLR7 is a key upstream driver of human SLE. Therapies blocking TLR7 itself or MyD88 may be more effective than therapies blocking GCs in patients with SLE due to increased TLR7 signalling.

**Online content**

Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at https://doi.org/10.1038/s41586-022-04642-z.
Article

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Methods

Mice
Mice were bred and maintained in specific-pathogen-free conditions at the Australian National University (ANU), Canberra, Australia. Experimentation was performed according to the regulations approved by the local institution ethics committee, including the Australian National University’s Animal and Human Experimentation Ethics Committee. Estimations of the expected change between experimental and control groups allowed the use of power analysis to estimate the group size that would enable detection of statistically significant differences. For in vitro experiments, randomization was not required given that there were no relevant covariates. Blinding was used for microscopy: histological analysis, electron microscopy imaging. Mice were used from 6–12 weeks, except for survival curves and tissue assessment (12–26 weeks). Both male and female mice were used and their genders are indicated in most figures (the Y chromosome is indicated in the genotype, that is, male mice).

Generation of the Tlr7- and Rnashe2b-mutant mouse strains
Tlr7<sup>−/−</sup> and deficient mice as well as Rnashe2b-deficient knockout mice were generated in a C57BL/6Ncrl background using CRISPR–Cas9-mediated gene editing technology. Genomic sequences were obtained from Ensembl (https://ensembl.org/) and compared to ascertain the conservation of the sequences between mouse and human genons. Single guide RNA (sgRNA) and single-stranded oligonucleotides were purchased from Integrated DNA Technology with the following sequences: Tlr7<sup>−/−</sup> sgRNA, 5′-TATGGACATTATACACTGGC-3′; Rnashe2b<sup>−/−</sup> sgRNA 5′-CTTTTAGTGCACTAGGTCAAGAACTTGCAACTCATTGAGGTTATTAAAATCATTTTCTTGGTATTTTCTTAAT-3′; and Rnaseh2b-R, 5′-AACACCTGCCCACATGTCAGGCAAATCCACTAGGTCACAATGCTAGGTAATGATATCAGTATAGACATTGAGGGAATTATTTCCACTTAGGTCAAGAACTTGCAACTCATTGAGGTTATTAAAATCATTTTCTTGGTATTTTCTTAAT-3′. The italicized nucleotides in the sgRNA sequences indicate the base altered by the respective variant in Tlr7 or Rnashe2b.

C57BL/6Ncrl female mice (aged 3–4 weeks) were mated with C57BL/6Ncrl males. Pseudopregnant CFW/crl mice were superovulated and mated with stud males. After detection of a vaginal plug, the fertilized zygotes were collected from the oviduct and Cas9 protein (50 ng µl<sup>−1</sup>) was co-injected with a mixture of sgRNA (2.5 ng µl<sup>−1</sup>) and single-stranded oligonucleotides (50 ng µl<sup>−1</sup>) into the pronucleus of the fertilized zygotes. After the micro-injection of the eggs, the zygotes were incubated overnight at 37 °C under 5% CO<sub>2</sub> and two-cell stage fertilized zygotes were collected from the oviduct and Cas9 protein was injected. Single-stranded oligonucleotides (50 ng µl<sup>−1</sup>) were incubated overnight at 37 °C under 5% CO<sub>2</sub> and two-cell stage fertilized zygotes were collected from the oviduct and Cas9 protein was injected. Single-stranded oligonucleotides (50 ng µl<sup>−1</sup>) were incubated overnight at 37 °C under 5% CO<sub>2</sub> and two-cell stage fertilized zygotes were collected from the oviduct and Cas9 protein was injected. For in vitro experiments, randomization was not required given that there were no relevant covariates. Blinding was used for microscopy: histological analysis, electron microscopy imaging. Mice were used from 6–12 weeks, except for survival curves and tissue assessment (12–26 weeks). Both male and female mice were used and their genders are indicated in most figures (the Y chromosome is indicated in the genotype, that is, male mice).

Flow cytometry
Single-cell suspensions were prepared from mouse spleens or thowed PBMCs, and individual subsets were analysed using flow cytometry. The primary antibodies used for mouse tissues included: SiglecH-APC (551, BioLegend), IgD–FITC (4057/18, BioLegend), IgD–PerCP Cy5.5 (HI-26.2a, BD Pharmingen), CD3–A700 (17A2, BioLegend), CD11b–PerCPCy5.5 (104, BD Bioscience), CD4–PerCPCy5.5 (RMA-4.5, BioLegend), IA/IE–BV421 (29F .1A12, BioLegend), CD138–PE (281-2, BD Pharmingen), PD1–BV421 (29F .1A12, BioLegend), CD19–BV510 (6D5, BioLegend), CD4–BV395 (6K1.5, BD Legend) CD21/35–BV605 (7G6, BD Horizon), CD45.1–BV605 (A20, BioLegend), CD45.1–BV711 (A20, BioLegend), CD45.1–PB (A20, BioLegend), Tlr7–PE (A94B10, BD Pharmingen), CD23–BV421 (B3B4, BioLegend), CXCR3–PE (CX3CR1-73, BioLegend), CD19–A700 (eBio1D3, Invitrogen), FOXP3–FITC (FKJ-16s, Invitrogen, eBioscience), FOXP3–PECy7 (FKJ-16s, Invitrogen, eBioscience), IgM–FITC/II (I/41, BD Pharmingen), IgM–PECy7 (II/41, Invitrogen), CD44–FITC (IM7, BD Pharmingen), CD44–PB (IM7, BioLegend), CD95 (FAS)–BV510 (Jo2, BD Horizon), BCL6–A647 (K12-91, BD Pharmingen), CD11b–PerCP Cy5.5 (M5/70, BioLegend), IA/IE–BV421 (M5/114.15.2, BioLegend), CD11c–A647 (N418, BioLegend), CD11c–BV510 (N418, BioLegend), CD11c–FITC (N418, BioLegend), CD25–PE (PC6, BioLegend), B220–A647 (RA3-6B2, BD Pharmingen), B220–BV395 (RA3-6B2, BD Horizon), B220–BV737 (RA3-6B2, BD Horizon), CD95–PECy7 (RI.388, BioLegend), CD4–PECy7 (RM4-5, BD Pharmingen), CD25–A647 (PC6, BioLegend), CD4–A647 (RM4-5, BioLegend), CD11c–APC (HL3, BD Bioscience), CD138–Biotin (281-2, BD Bioscience), CXCR5–Biotin (2G8, BD Bioscience), streptavidin-BV805 (BD Horizon), streptavidin-BV510 (BD Biosciences, Invitrogen), Streptavidin-PECy7 (BD Horizon), streptavidin-A700 (BioLegend), Streptavidin-Cy5 (BD Biosciences). For human PBMCs: CD19–BV605 (BD Biosciences, Invitrogen), Streptavidin-Cy5 (BD Biosciences). For human PBMCs of TLR7 and Rnashe2b mutant mouse strains.

For human PBMCs: CD19–BV605 (BD Biosciences, Invitrogen), Streptavidin-Cy5 (BD Biosciences). For human PBMCs of TLR7 and Rnashe2b mutant mouse strains.
Fc receptors were blocked using purified rat anti-mouse CD16/CD32 (Mouse BD Fc Block, BD Biosciences) and then stained for 30 min at 4 °C in the dark, with primary and secondary antibodies. Intracellular staining was performed using the FOXP3 Transcription Factor Staining Buffer Set (eBioscience) according to the manufacturer’s instructions. Samples were acquired on the Fortessa or Fortessa X20 cytometer with FACSDiva (BD, Biosciences) and analysed using Flowjo v.10 (Flowjo). All fluorescence-activated cell sorting (FACS) and microscopy analysis was carried out at the Microscopy and Cytometry Facility, Australian National University.

Sanger sequencing
Primers for human TLR7 DNA sequencing were used at 10 μM (primer sequences available on request). PCR amplification was carried out using Phusion Hot Start II DNA Polymerase II (Thermo Fisher Scientific) and under the conditions recommended by the manufacturer. PCR amplicons were electrophoresed and excised bands were purified using the QIAquick Gel Extraction Kit (Qiagen). Sanger sequencing was completed using Big Dye Terminator Cycle sequencing kit v3.1 (Applied Biosystems) using the same primers used for PCR amplification. Sequencing reactions were run on the 3730 DNA Analyzer (Applied Biosystems) system at the ACRF Biomolecular Resource Facility, Australian National University.

Immunohistochemistry
Liver, pancreas and kidneys were fixed in 10% neutral buffer formalin solution, embedded in paraffin and stained with H&E.

Bone marrow chimera experimentation
For competitive bone marrow chimeras, Rag1−/− mice were irradiated and injected intravenously with equal numbers of bone marrow cells from either wild-type or kika CD45.2 and wild-type CD45.1 mice. Mice and injected intravenously with equal numbers of bone marrow cells from either wild-type or kika CD45.2 and wild-type CD45.1 mice. Mice were given Bacitracin in their drinking water for 48 h before injection and for 6 weeks after injection, and housed in sterile cages. After 22 weeks of reconstitution, mice were taken down for phenotyping by flow cytometry.

B cell culture and Cell Trace Violet staining
Single-cell suspensions were prepared from kika, wild-type or Tlr7kik/Y mice and graphed using PRISM.

BMDM culture and stimulation
Primary BMDMs from 3 Tlr7+/+ mice and wild-type littermates were extracted and differentiated for 7 days in complete DMEM supplemented with L929-conditioned medium as previously reported, before overnight stimulation with ssRNA, guanosine or R848. Noticeably, the yield of Tlr7+/+ BMDMs obtained after 7 day differentiation was substantially greater than from wild-type mice. All synthetic RNAs were synthesized by Integrated DNA Technologies. ssRNAs (below) with no backbone modification were resuspended in duplex buffer (100 mM potassium acetate, 30 mM HEPES, pH 7.5, DNase-RNase-free H2O), and were previously shown to induce TLR7 sensing in human cells48, ssRNAs were transfected with DOTAP (Roche) and pure DMEM in biological triplicate, as previously described48, to a final concentration of 500 nM. The ratio of DOTAP to RNA (at 80 μM) was 3.52 μg μl−1 of ssRNA. Guanosine (Sigma-Aldrich, G6264, 10 mg freshly resuspended in 176.5 μl DMSO (200 μM stock solution)) and R848 (Invivogen, tlr848) were used at the indicated final concentrations. TNF levels in culture supernatants were detected using the BD OptEIA Mouse ELISA kit (BD Biosciences) according to the manufacturers’ protocols. Tetramethylbenzidine substrate (Thermo Fisher Scientific) was used for quantification of the cytokines on a Fluostar OPTIMA (BMG LABTECH) plate-reader. The RNA sequences used (5′-3′) were as follows: B-406AS-1, GCCGGACAUUAUUUAUACGC; 41-6, GCCGGACAUAUUUAUACGC; 41-8, GCCGGUCUUAAUUAUACGC; 41-10, GCCGCUUUCUUUAAUCGC.

ADVIA blood analysis
Orbital bleeds were performed on mice and blood samples were run on the ADVIA system (Siemens Advia 1200).

Western blotting
Cytosolic extracts were prepared from around 20 million–40 million splenocytes by lysis in Triton X-100 buffer (0.5% Triton X-100, 20 mM Tris- HCl pH 7.4, 150 mM NaCl, 1 mM EDTA, 10% glycerol) and centrifuged. Cytosolic extracts were resolved on 8% SDS–polyacrylamide gels and probed with the relevant primary and secondary antibodies. Rabbit anti-TLR7 (D7; Cell Signaling Technology) and mouse anti-mouse TLR7-PE (A94B10; BD Biosciences) were used at 1:1,000, the actin monoclonal antibody (JLA20, Developmental Studies Hybridoma Bank, The University of Iowa) was used at 1:5,000. Membranes were developed with Clarity Western ECL Substrate (BioRad Laboratories).

Dual-luciferase assays
RAW264.7 cells were transfected with 245 ng of pNIFTY (NF-κB lucerase; InvivoGen), pRL-CMV (100 ng, Promega) Renilla lucerase control plasmid, 125 ng of TLR7-RA plasmids (Genecopoeia) expressing the individual variants. After overnight expression, half of the samples were stimulated with 1 mM 2′,3′-cGMP (Santa Cruz) or 1 mM guanosine plus 20 μg ml−1 ssRNA using DOTAP for 6 h and dual-luciferase assays were performed as previously described43. Raw264.7 cells (originally from ATCC) were tested for mycoplasma contamination using Plasm Test (InvivoGen).

Statistics
Statistical analysis was carried out using R software v.3.6.1 (The R Foundation for Statistical Computing) and the Emmeans package. Mouse spleen mass data were analysed using two experiments as a blocking factor and one-way ANOVA, followed by a pairwise estimated marginal means comparison of genotypes. Mouse cellular phenotyping, ELISAs, white blood cell and platelet count analyses were performed using a log linear regression model and one-way ANOVA, followed by a pairwise estimated marginal means comparison of genotypes. Purified B cell cultures were analysed using a linear regression model and one-way ANOVA, followed by a pairwise estimated marginal means comparison of genotypes and stimulatory effect. Lucerase assay statistics were analysed using one-way ANOVA with Bonferroni multiple-comparison test (Prism, GraphPad). All data were filed using Microsoft Excel 2016 and graphed using PRISM.
DNA, RNA and nRNP ELISAs
Plates were coated with poly-L-lysine (Sigma-Aldrich) before addition of 2.5 μg of either DNA (D7290, Sigma-Aldrich), RNA (AM7120G, Thermo Fisher Scientific) or nRNP (SRC-1000, Immunovision). Plates were then blocked in ELISA blocking buffer (PBS and 1% BSA) for 2 h at room temperature. Mouse serum was diluted 1:40 with ELISA coating buffer (0.05 M sodium carbonate anhydrous/sodium hydrogen carbonate, pH 9.6), and incubated in the ELISA plates overnight at 4 °C. The plates were washed and goat anti-mouse IgG-AP antibodies (Alkaline Phosphatase, Southern Biotech) were added for 1 h at 37 °C. Phosphatase substrate (Sigma-Aldrich, S0942) was used as described by the manufacturer. The samples were read using the Infinite 200 PRO Tecan Microplate Reader (Tecan Group) at an absorbance of 405 nm and normalized to background absorbance at 605 nm.

Hep-2/C. luciliae immunofluorescence
ANAs and dsDNA were determined using Hep-2 and Crithidia luciliae slides (both from NOVA Lite, respectively). Serum was diluted 1:40 for Hep-2 slides and 1:20 for Crithidia slides and stained as described by the manufacturer using donkey anti-mouse IgG Alexa-488 (Molecular Probes) as the secondary antibody. The slides were imaged using an Olympus IX71 inverted fluorescence microscope.

RNA-seq analysis
Total B cells were obtained from wild-type or kika mouse spleens and purified using the Mouse B Cell Isolation Kit (Miltenyi Biotec) and stimulated with anti-mouse IgM (10 μg ml−1) for 20 h. Total RNA was extracted using RNEasy Mini Kits (74104, Qiagen). Sequencing was performed using the NextSeq500 platform and analysis was conducted using the following R packages: limma, edgeR and enhanced volcano. For the patient, type I IFN single-cell RNA-seq analysis was performed. PBMCs were isolated from frozen human samples as previously described. Live cells were next purified by FACS using 7AAD and labelled with TotalSeq anti-human-histag (BioLegend). The number of cells was determined and 10,000 cells per sample were run on the 10x Chromium platform (10x Genomics). Library preparation and sequencing were performed by The Biomedical Research Facility according to the manufacturer’s instructions for the Chromium Next Gem Single Cell 5’ Kit v2. The samples were sequenced using the NovaSeq 6000 (Illumina) system. The FASTQ files were aligned to the human GRCh38 reference genome using 10x Genomics Cell Ranger pipeline v6.0.1. Statistical analysis, clustering and visualization were conducted using Seurat v4.0.1 in the R environment.

Molecular dynamics simulations
Details of the computational modelling are provided in the Supplementary Methods.

Reporting summary
Further information on research design is available in the Nature Research Reporting Summary linked to this paper.

Data availability
For the genomic data relating to the families in Fig. 1, family A has been deposited in SRA, under BioProject accession number PRJNA798834; family B has been deposited at the Baylor-Hopkins Center for Mendelian Genomics, AnVIL repository under the participant IDs BH14435-1 (proband), BH14435-2 (mother) and BH14435-3 (father); family C has been submitted to the EGA (EGAS00001005965). Sequencing data in Fig. 3i have been deposited at the Gene Expression Omnibus under accession number GSE196316 and Fig. 4o has been deposited at Open Science Framework (OSF) under accession number pW6zn.

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Extended Data Fig. 1 | Confirmation of paternity in trio of proband with TLR7 de novo variant and methods for molecular modelling. (a) Peddy diagrams used to establish relatedness. Each red dot represents a child/parent pair (child mother and child father). The grey dot is a no-relatedness control. Coefficient of relatedness should be 0.5 for a parent-child pair. ibs0: the number of sites at which the 2 samples shared no alleles (should approach 0 for parent-child pairs). ibs2: the number of sites in which the child vs parent samples were both hom-ref, both het, or both hom-alt. Shared_hets: the number of sites at which both child and parent samples were hets. (b) Ancestry check using Peddy (proband and parents are purple dots). (c) Phylogenetic conservation of TLR7 variants. (d) Integrative Genomics Viewer (IGV) image of the Y264H TLR7 de novo variant. (e) TLR7 structure 6IF5. Regions in red were restrained through all simulations with a harmonic restraint of force constant 5 kcal/mol/Å², and correspond to residue numbers: 27-96, 116-179, 193-256, 281-297, 304-346, 361-376, 385-403, 412-427, 434-460, 476-499, 510-523, 534-548, 561-572, 591-602, 616-625, 646-656, 671-681 and 699-835. (f) Guanosine and R848 illustrated with binding geometries from crystal structures 5GMF and 5GMH14. L1-L3 indicate ligand atoms used for Boresch restraints, which were restrained relative to the three depicted protein alpha carbons of residues F408, G379 and F325 (not to scale). Distances and angles in gold, and dihedrals in pink show the values for the 6DoF Boresch restraints. Additional geometric relationships between the restrained atoms, as measured from the starting structure, are shown in grey smaller print. Boresch dihedral restraints are relative to the two atoms connecting either side of the location of print. White hydrogen spheres and red oxygen spheres show the atoms used in the calculation for determining the number of waters within 3.5 Å of the tail region that each ligand interacted with.
Extended Data Fig. 2. Tlr7−/− spleen cells lack TLR7 expression. (a) Tlr7 nucleotide and amino acid sequence in mice carrying a CRISPR/Cas9-generated deletion (Tlr7−/−) and WT littermate (Tlr7+/+). (b) Flow cytometric histograms of intracellular TLR7 expression on cells from 6-month-old mice of the indicated genotypes: plasmacytoid dendritic cells (pDC, CD11c−CD19−SiglecH+; B cells (CD19+) (EPMF)). Bars represent medians and each dot a single mouse. These results are representative of one experiment. One-way ANOVA with Tukey (b); Exact p values are shown.
Extended Data Fig. 3 | Autoantibodies to ssDNA but not dsDNA are a feature of kika mice and are detectable by 6 wks of age. (a) Quantification of ANAs in 12 wk-old kika mice by Hep-2 immunofluorescence (IF). (b) Proportion of 6 month-old WT and kika mice positive for dsDNA according to Crithidia luciliae IF, and representative images (c) for each genotype. (d) Autoantibodies to DNA in serum from 4 wk-old (n = 10) and 6 wk-old (n = 9) wt or kika mice. (e) HEp-2 IF showing pattern and quantification of ANAs in 6 wk-old kika mice. (f) Mesoscale measurement of cytokines in serum from wt or kika mice (n = 20). (g) Dose-dependent response of TLR7 to guanosine (data averaged from two mice in biological triplicate). (h) Responsiveness of TLR7 to ssRNAs lacking uridine (ss41-L), with 6 to 10 uridines; or 9 uridines B-406-AS1 (data represent mean ± s.e.m. averaged from three mice in biological triplicates). Unpaired t-test (d, h); Mann-Whitney test (f). Exact p values are shown.
Extended Data Fig. 4 | Organ Pathology of kika mice. Observational report is summarized in supplementary table 4. Organs were collected from 3 mice per genotype. Histopathology and organ pathology was performed by the Australian Phenomics Network (APN).
Extended Data Fig. 5 | See next page for caption.
Extended Data Fig. 5 | Cellular phenotypes in blood and spleen from kika mice, TLR7-deficient mice, mixed chimeras and Yaa mice. (a) White blood cell (WBC) count in 18-wk old mice. (b, c) Flow cytometric plots and quantification. (b) Spleen T (CD3+):B (B220+) cell ratio from 12-wk old kika mice. (c) Age-associated B cells (ABC, B220+ CD21- CD23- CD19hi CD11c+) in blood from 18-wk-old kika mice. (d) Splenic marginal zone (MZ) B cells (CD19+ CD23- CD21+) in 12-wk kika mice. (e) Circulating T follicular cells (Tfo, CD4+ CXCR5+ PD1hi) and extrafollicular helper T cells (eTf, CD4+ CXCR5+ PD1+ CXCR3+) in blood from 18-wk-old kika mice. (f) Plasmacytoid dendritic cell (pDCs, CD3- CD19- MHCII+ CD11c+ CD11b- CD8- SiglecH+ BST2+) and pDC MFI of MHCII and SiglecH from 12-week-old kika mice. (g) Splenic germinal center B cells (GCB, CD19+ CD95+ BCL6+), ABC (B220- CD21- CXCR5- CD19hi CD11c+), Tfo (CD4+ CXCR5+ PD1hi), eTfhi (CD4+ CXCR5+ PD1+ CXCR3+) and plasma cells (PC, CD138+ CD98-) from 24-wk TLR7-deficient mice. (h, i) Splenic cell subsets from mixed bone marrow chimeric mice containing a 1:1 ratio of control Tlr7+/+ CD45.1/ Tlr7+/+ CD45.2 or Tlr7+/+ CD45.1/Tlr7kik/kik CD45.2 bone marrow (h) and 100% mixed bone marrow (i) of each genotype. Subsets shown are CD4 effector (CD4+ FoxP3- CD44+), MZ (CD19+ CD23- CD21+), Treg (CD4+ FoxP3+) and CD45.1 to CD45.2 reconstitution ratio, 22-weeks post-reconstitution. (j) Autoantibodies to DNA and smRNP in serum from 100% chimeric mice model. (k) Splenic cell subsets from kika and Yaa mice. Bars represent medians and each dot a single mouse. These results are representative of one blood ADVIA analysis, two experiments for blood flow cytometry, four splenic phenotyping for kika mice and one for TLR7 deletion mice, and one experiment for chimera analysis. One-way ANOVA with Tukey (a–g, k); Two-way ANOVA (h); Mann-Whitney (i); Unpaired t-test (j); Exact p values are shown.
Extended Data Fig. 6 | See next page for caption.
Extended Data Fig. 6 | Rnaseh2b hemizygosity does not cause a cellular phenotype. (a) Rnaseh2b cDNA sequence from Rnaseh2b-deletion mice, highlighting the single nucleotide CRISPR/Cas9-generated deletion leading to a frame-shift after amino acid residue Q170 and stop codon 4 amino acids downstream. (b, c) Flow cytometric quantification of splenic germinal center B cells (GCB, CD19+ CD95+ BCL6+), age-associated B cells (ABC, B220+ CD21+ CXCR5- CD19+ CD11c-), T follicular helper cells (TFH, CD4+ CXCR5+ PD1hi), extrafollicular helper T cells (eTH, CD4+ CXCR5- PD1+ CXCR3+) and plasma cells (PC, CD138+ CD98+) (b) in 12-week-old mice carrying a heterozygous deletion in Rnaseh2b and (c) in 12-wk-old mice with heterozygous deletion in Rnaseh2b crossed to the kika mice. (d) Autoantibodies to DNA, RNA and smRNP in serum from mice of indicated genotypes. (e) Representative photo of time-mated embryos from Rnaseh2b+− breeders. (f) Western blot of RNASEH2B in time-mated embryo lysates. (g) KASP genotyping results of representative time-mated embryos. (h) Breeding record of Rnaseh2b+− crossed to Rnaseh2b+− mice. (i) Type I IFN signature of Rnaseh2b+/− Tlr7+/−/− double heterozygous female mice compared to Tlr7+/−/− alone, Rnaseh2b+/−/− alone or Trex1−/− mice as positive controls. Bars represent medians and each dot a single mouse. Data is representative of two experiments. Embryos were collected from two time-mating breeding set ups (n = 18 embryo collected from two pair breeders). One-way ANOVA with Tukey (b–d); Chi-square test (h); Exact p values are shown.
Extended Data Fig. 7 | *Kika* mice have normal responses to TLR7 signalling and CD93 is expressed on TLR7 stimulated B cells. (a–c) Flow cytometric analysis of (a) Mean CD93+ cells derived from CD93- sorted splenic B cells stimulated with R837, α-IgM or R837 + α-IgM for 72-hours from *kika* and control mice. (b) FACS sorted bone marrow immature B cells (*B220^int^CD93^+^) cultured with or without α-IgM for 72 h, from male mice of the indicated genotypes. (c) Mean survival count of MACS purified splenic B cells stimulated with R837, α-IgM or R837 + α-IgM for 72-hours from *kika* mice and control mice. (c) Percentage of bone marrow (BM) immature B cells (CD93^+^ B220^+^) in 12-35 wk *kika* and control mice. (a, b, d) Bars represent means ± s.d. and (d) each dot a single mouse. These results are representative of one experiment for CD93 upregulation and BM immature flow cytometry, four splenic B cell cultures purified using MACS bead selection and one BM analysis. Two-way ANOVA with Tukey (a, b) and Sidak (d); Exact p values are shown.
Extended Data Fig. 8 | Myd88 deficiency rescues kika's immune cell phenotype and splenic cellularity. (a) Flow cytometric quantification of splenic cell subsets (percentage and total number) and total cellularity in 12 wk-old mice of the indicated genotypes (Tlr7+/+Myd88+/+ n = 12, Tlr7+/+Myd88−/− n = 11, Tlr7+/+Myd88+/− n = 4). Subsets include: germinal center B cells (GCB, CD19+ CD95+ BCL6+), T follicular helper cells (TFH, CD4+ CXCR5+ PD1hi), percentage of ABC (B220+, CD21−, CXCR5− CD19hi CD11c+), extrafollicular helper cells (eT H, CD4+ CXCR5− PD1+ CXCR3+), plasma cells (PC, CD138+ CD98+), CD4 effector/memory T cells (CD4+ FoxP3− CD44+ CXCR5+ PD1hi), and splenic cellularity (% of CD4+ and # of CD4+). Male mice = grey and female mice = white. Bars represent medians and each dot a single mouse. One-way ANOVA with Tukey; Exact p values are shown.
Extended Data Fig. 9 | TLR7 and MYD88 expression is indistinguishable between female patients with SLE and healthy controls (HC). (a) Flow cytometric analysis and quantification of DN B cells, pDCs and ABCs phenotype in PBMCs from healthy controls (n = 6) and female patients with SLE (n = 8). (b) Quantification of TLR7 and MYD88 protein in HC and SLE. (c) Quantification of CD25 expression after TLR7 stimulation among HC (n = 8), autoimmune control (AC) (n = 1), age-matched autoimmune control (AMAC) (n = 1) and A.II.1 (n = 1). Unpaired t-test (a, b); One-way ANOVA with Tukey (c); Exact p values are shown.
Reporting Summary

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Statistics

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

- The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
- A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
- The statistical test(s) used AND whether they are one- or two-sided
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- For null hypothesis testing, the test statistic (e.g. F, t, χ²) with confidence intervals, effect sizes, degrees of freedom and P value noted. Give P values as exact values whenever suitable.
- For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
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- Estimates of effect sizes (e.g. Cohen’s d, Pearson’s r), indicating how they were calculated

Our web collection on statistics for biologists contains articles on many of the points above.

Software and code

Policy information about availability of computer code

Data collection
Cellular phenotype data was collected on a Fortessa or Fortessa X-20 cytometer running FACSDiva version 8.0 (BD, Biosciences).

Data analysis
Statistical analysis was carried out using R software version 3.6.1 (The R Foundation for Statistical Computing) and the Emmeans package. Luciferase assays statistics were analysed using a one-way ANOVA with Tukey’s multiple comparison (PRISM 6, GraphPad Software LLC). Flow cytometry data was analysed using the FlowJo software v10 (FlowJo LLC). Microsoft Excel 2016, Sequencher v5. Single-cell data was analysed using Cell Ranger v6.0.1 pipeline.

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.

Data

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:
- Accession codes, unique identifiers, or web links for publicly available datasets
- A list of figures that have associated raw data
- A description of any restrictions on data availability

The authors declare that data supporting the findings of this study are available within the paper or its supplementary information files. Data that support the findings of this study are also available from the corresponding author (CSV) upon reasonable request.
**Field-specific reporting**

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## Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

| Sample size | For each experiment we estimated the expected change between experimental and control groups (e.g. at least a 20% change and SD at most half the magnitude of the minimum effect size we were interested in). With these assumptions we used power analysis to estimate the group size that would provide at least 80% power to detect statistically significant difference (with p < 0.05 considered significant). |
|-------------|---------------------------------------------------------------------------------------------------------------|
| Data exclusions | No data was excluded |
| Replication | Figs 1a-d, g-i, 1k (survival curve) no experimental replication is feasible.  
NF-kB luciferase assays in Fig 1e, f is representative of 2 experiments.  
Fig 1j results are representative of 4 experiments for spleen mass and 4 pooled experiments for cellularity, 3 for serum ELISAs (Fig 1m), 1 times for ANAs (Fig 3l).  
Figure 1n (R848) used 2 mice per genotype (done in triplicate wells) and done on the same day. Fig 1o (Guanosine) used 3 mice per genotype (all in triplicate wells). Done on two different days.  
Platelet analysis (Fig 2a) was done once and the organ H&E analysis and electron microscopy (Fig 2b-c) 3 times. H&E stain in fig 2d was done once, and serum level in Fig 2e were assayed once using large numbers of mice. Figs. 2f-h,j,k of the splenic phenotyping in Figs 2f-h,j,k are representative of four experiments whilst the kidney phenotyping analysis (Fig 2j) is representative of two experiments.  
Fig 3a-b were done once. Fig 3g, the 8 cell culture was done 3 times.  
Fig 3c-d: representative of two experiments, and Fig 3h is a compilation of two experiments. No experimental replication was feasible for Fig 3i. Western blotting results in Fig 3e-f are representative of 3 experiments each.  
Figs 4a-e, the MyD88/-/- crossovers were done once. Figs. 4f-h are representative of two experiments and the Western blots in Fig 4i were replicated at least 2 times for each antibody. Figs. 4j-o: no experimental replication was feasible.  
Fig 5a-f no experimental replication feasible. Fig 5a-b were done once.  
Fig 5a, b, c was done once.  
Fig 5d-f is representative of 1 experiment. Fig 5g (dose response) used 2 mice per genotype (all in triplicate wells). Done on the same day.  
Fig 5h (ssRNA): used 3 mice per genotype (all in triplicate wells). Done on two different days.  
Fig 5c was carried out once on organs from 3 mice of each genotype.  
Fig 5e-f: adenovirus blood analysis and analyses of chimeras in Figs 5h-j were done once, blood flow cytometry twice (Fig 5c, e) and splenic flow four times for Fig 5b, f, g and h. Fig 5i is a compilation of 2 experiments.  
Fig 5b-g results are representative of 2 experiments. Fig 5h is a compilation of five litters of genotyping results. Fig 5i was done twice.  
Fig 5a, c, d is representative of 1 experiment, whilst 4 splenic B cell cultures were used in Fig 5j.  
Fig 5a-b was done once.  
Fig 5a-c was done once. |
| Randomization | For in vitro experiments, randomisation was not required given there were no relevant covariates (i.e. cells from littermate mice came from the same cage, all wells treated simultaneously using multi-channel pipettes, on the same day, in the same single plate, analysed in the same machine, handled by the same investigator). |
| Blinding | Blinding to allocation occurred for all experiments in which the investigator had to score data manually (i.e. intensity and pattern of ANA fluorescence, analysis of histological samples from mouse necropsies, assessment of Ig deposits in EM kidney sections). Blinding did not occur for assays in which a pre-determined order was required for loading gels plus the result would be presented raw to the reader (i.e. western blot gels) or analysed via an automated machine without input from the investigator (i.e. quantification of luciferase activity). Investigators planned mouse experiments based on genotype and grouping, but during performance of experiments mice were identified only by randomly assigned number with investigators blind to group allocation. |

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## Reporting for specific materials, systems, and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.
## Materials & experimental systems

| n/a | Involved in the study |
|-----|-----------------------|
| ☒  | Antibodies            |
| ☒  | Eukaryotic cell lines |
| ☒  | Palaeontology and archaeology |
| ☒  | Animals and other organisms |
| ☒  | Human research participants |
| ☒  | Clinical data |
| ☒  | Dual use research of concern |

## Methods

| n/a | Involved in the study |
|-----|-----------------------|
| ☒  | ChiP-seq |
| ☒  | Flow cytometry |
| ☒  | MRI-based neuroimaging |

### Antibodies

**Antibodies used** All antibodies used are commercially available and extensively used. We have listed all antibodies and their clone names in the materials section but given the large number of antibodies used over the breath of the work we did not note all their lot numbers. SigleC1-APC (#551, Biologend), IgD-FTC (#405718, Biologend), IgD-PerCP Cy5.5 (#11-26c.2a, BD Pharmingen), CD3-ALTO (#17A2, Biologend), CD19- BV395 (#1D3, BD Horizon), CD138-PE (#281-2, BD Pharmingen), PDI-BV421 (#29F.1A12, Biologend), CCR7-PerCP Cy5.5 (#4812, Biologend), CD8-BV405 (#53-6.7, BD Horizon), CD19-BV510 (#605, Biologend), CD4-BV395 (#66K1.5, BD Horizon) CD213J5-BV605 (#76G, BD Horizon), CD45.1-BV605 (#20A, Biologend), CD45.1-BV711 (#2A3, Biologend), CD45.1-BV605 (#2A0, Biologend), CD45.1-BV605 (#2A0, Biologend), CD19-PE (#44G2, BD Pharmingen), CD127-PE (#450, BD Pharmingen), CD25-PE (#8A3, BD Pharmingen), CD86-PE (#55F5, BD Pharmingen), CD28-PE (#73.2A3, BD Pharmingen), CD69-PE (#F5H2, BD Pharmingen), HLA-DQ-PE (#65302, Biologend), HLA-DR-PE (#24-225, BD Pharmingen), CD8-PE (#53-8, BD Pharmingen), CD5-PE (#573A11, BD Pharmingen), CD4-PE (#10F.1, BD Pharmingen), CD3-PE (#17A2, BD Pharmingen), CD25-PE (#2B11, BD Pharmingen), CD25-PE (#3C7, BD Pharmingen). All antibodies were used as recommended by the manufacturers. All antibodies were used and had been previously validated by the manufacturing companies. We further validated mouse TLR7-PE A94B10 antibody by staining splenocytes from TLR7 knockout mice. We provide the clones used for each antibody. Antibody titrations and dilutions used in each experiment are only relevant to the specific batch used, which change over time and therefore not useful.

### Eukaryotic cell lines

**Policy information about cell lines**

| Cell line source(s) | RAW264.7 cells were originally from the American Type Culture Collection (ATCC) |

**Authentication**

The cell line used has not been authenticated by STR profiling.

**Mycoplasma contamination**

All cell lines tested negative for mycoplasma contamination using PlasmoTestTM (InvivoGen).

**Commonly misidentified lines**

(See ICLAC register)

The cell line used is not listed in the database of commonly misidentified cell lines.

### Animals and other organisms

**Policy information about studies involving animals; ARRIVE guidelines** recommended for reporting animal research

| Laboratory animals | C57BL/6 Ncr1 mice were used in this study. Both male and female mice were used and the sex has been identified in all figures by the genotype, adding the "Y" in all male genotypes, and specifying the allele of the X chromosomes; (+ or kik). Mice were used at 8-12 weeks for phenotyping and in vitro experiments, except organ histology which was examined at 26 weeks. |
CFW/crl female mice from 6-16 weeks of age were used to mate with stud males for generation of CRISPR/Cas9 Tlr7 and Rnaseh2b gene edited mice.

| Wild animals                      | The study did not involve wild animals. |
|-----------------------------------|----------------------------------------|
| Field-collected samples           | The study did not involve field animals.|
| Ethics oversight                  | Animal experimentation was performed according to the regulations approved by the Australian National University's Animal Experimentation Ethics Committee. |

Note that full information on the approval of the study protocol must also be provided in the manuscript.

**Human research participants**

Policy information about **studies involving human research participants**

**Population characteristics** Individuals were either healthy controls, or patients who were diagnosed with systemic lupus erythematosus by treating physicians, or their family members. Individuals known medical treatments and clinical diagnosis are provided in Table S1.

**Recruitment** Participants were recruited by their referring medical practitioners, based on clinical eligibility criteria, with the vast majority of eligible participants agreeing to participate in the study. Young healthy controls for the phenotyping/RNAseq experiments of the proband with the TLR7 Y264H variant were recruited amongst young teenage girls within the same school in Canberra, of white European ascent (including one Spanish); thus gender, age, and ethnically matched to the proband (Spanish). However, other environmental/geographic influences (proband had been living in Guatemala for several years when last bled) could not be controlled for.

| Ethics oversight | The study was approved by and complies with all relevant ethical regulations of the Australian National University and ACT Health Human Ethics Committees (2015/079), the University Hospitals Institutional Review Board, or by Renji Hospital Ethics Committee of Shanghai Jiaotong University School of Medicine. |

Note that full information on the approval of the study protocol must also be provided in the manuscript.

**Flow Cytometry**

**Plots**
- Confirm that:
  - The axis labels state the marker and fluorochrome used (e.g., CD4-FITC).
  - The axis scales are clearly visible. Include numbers along axes only for bottom left plot of group (a ‘group’ is an analysis of identical markers).
  - All plots are contour plots with outliers or pseudocolor plots.
  - A numerical value for number of cells or percentage (with statistics) is provided.

**Methodology**

**Sample preparation** Human PBMCs were isolated using Ficoll-Paque gradient centrifugation and frozen thawed before staining for flow cytometric analysis. Single cell suspensions were prepared from mouse spleens and B cells were magnetically purified using mouse B Cell Isolation Kit (Miltenyi Biotec), labeled with Cell Trace Violet (CTV, Thermo Fisher) and cultured for 72 hours in complete RPMI 1640 media (Sigma-Aldrich) supplemented with 2mM L-Glutamine (GIBCO), 100 U penicillin-streptomycin (GIBCO), 0.1 mM nonessential amino acids (GIBCO), 100 mM HEPES (GIBCO), 55 mM β-mercaptoethanol (GIBCO) and 10% FBS (GIBCO) at 37°C in 5% CO2. Bone marrow was obtained from mice, the Fc receptors blocked and cells stained and sorted.

**Instrument** Cells were sorted on a FACS Aria II, splenocytes and human PBMC samples were acquired on a Fortessa or Fortessa X-20 cytometer.

**Software** FACS data was analyzed using FlowJo software v10 (FlowJo LLC).

**Cell population abundance** Sorted sample purity was based on flow cytometry sorting analysis and stringent gating. Abundance of populations are indicated in the gating figures of the manuscript.

**Gating strategy** SC-H/FSC-A (cells were gated along a diagonal gating strategy to eliminate cells with disproportional FSC-H and FSC-A size), SSC-W/SSC-H (cells with large SSC-W from scatter were eliminated), FSC-A/live dead (cells staining negative for the live dead marker were selected as "live") and FSC-A/SSC-A (Cells were gated as lymphocytes if they had a lower size and granularity relative to other signals detected). Once cells were established as singlets, live and lymphocytes analysis was completed as described in the manuscript, where possible biphasic populations were used to identify positive and negative populations. The gating strategies used to identify individual subsets of interest have all been shown in the main figures.

Tick this box to confirm that a figure exemplifying the gating strategy is provided in the Supplementary Information.