

## ORIGINAL ARTICLE

*Irf4 is a positional and functional candidate gene for the control of serum IgM levels in the mouse*J Côrte-Real, J Rodo, P Almeida, A Coutinho, J Demengeot and C Penha-Gonçalves  
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Natural IgM are involved in numerous immunological functions but the genetic factors that control the homeostasis of its secretion and upholding remain unknown. Prompted by the finding that C57BL/6 mice had significantly lower serum levels of IgM when compared with BALB/c mice, we performed a genome-wide screen and found that the level of serum IgM was controlled by a QTL on chromosome 13 reaching the highest level of association at marker D13Mit266 (LOD score = 3.54). This locus was named IgMSC1 and covered a region encompassing the interferon-regulatory factor 4 gene (*Irf4*). The number of splenic mature B cells in C57BL/6 did not differ from BALB/c mice but we found that low serum levels of IgM in C57BL/6 mice correlated with lower frequency of IgM-secreting cells in the spleen and in the peritoneal cavity. These results suggested that C57BL/6 mice have lower efficiency in late B-cell maturation, a process that is highly impaired in *Irf4* knockout mice. In fact, we also found reduced *Irf4* gene expression in B cells of C57BL/6 mice. Thus, we propose *Irf4* as a candidate for the IgMSC1 locus, which controls IgM homeostatic levels at the level of B-cell terminal differentiation. Genes and Immunity (2008) 0, 000–000. doi:10.1038/gene.2008.73

**Keywords:** serum IgM; homeostasis; IRF4**Introduction**

Natural IgM play a significant role in the immunology of health and disease. IgM antibodies are involved in numerous immunological functions including protection against infection,<sup>1–3</sup> prevention of autoimmunity,<sup>4,5</sup> B-cell homeostasis<sup>6–8</sup> and immunosurveillance against tumors.<sup>9,10</sup> IgM biological roles also include opsonization of apoptotic cells, resulting in accelerated clearance by phagocytic cells.<sup>11</sup> Many of the circulating IgM correspond to natural antibodies that exist prior to infection or immunization.<sup>12–14</sup> Natural IgM are polyreactive to evolutionary conserved structures<sup>15</sup> and bind efficiently to previously un-encountered antigen.<sup>16</sup> These antibodies compensate their low-antigen affinity with relatively high avidity and furthermore the effectiveness of the antigen–antibody interaction is enhanced by the high efficiency of IgM in engaging the complement pathway.<sup>11,17</sup>

The preimmune Ig repertoire is thought to be composed by mature B cells either recirculating through follicles of secondary lymphoid organs (B2 cells), or joining compartments in specific locations as the marginal zone in the spleen (MZ B-cells) and the pleural or peritoneal cavities (B1 cells).<sup>12</sup> B2 and marginal zone B-cell development is initiated in the bone marrow and completed in the periphery<sup>18,19</sup> throughout life. B cells

exported from the bone marrow mature in the spleen<sup>20</sup> into subsets that differ in their surface phenotype, anatomic localization and immunologic function<sup>21–23</sup> and that ultimately are able to differentiate into antibody-producing plasma cells, upon antigenic stimulation.<sup>18</sup> Most frequently primary antigen contact of mature naive B cells leads to IgM production detectable in the serum.

On the other hand, studies on germ-free and axenic mice suggest that in the absence of pathogens such stimulation is conveyed by the action of self-antigens, leading to production of natural antibodies.<sup>24,25</sup> The B1 cell population, is derived almost exclusively from foetal liver,<sup>26</sup> is established early in ontogeny and is predominantly maintained by self-replenishment.<sup>27</sup> These cells locate preferentially in the pleural and peritoneal cavities<sup>26</sup> and spontaneously secrete natural Ig<sup>28</sup> participating in relatively few antigen-stimulated antibody responses.<sup>26</sup>

The capacity for Ig secretion is acquired in the process of B-cell terminal differentiation,<sup>29</sup> at the plasma cell stage and requires the expression of the transcription factor *Irf4*,<sup>30</sup> a member of the interferon-regulatory factor (IRF) family.<sup>31</sup> *Irf4* expression is induced by antigen receptor or mitogen stimulation<sup>32</sup> and is restricted to cells of the immune system,<sup>33,34</sup> playing a critical role in both early<sup>35</sup> and late B-cell differentiation.<sup>30</sup> Interestingly, *Irf4* upregulation results in spontaneous differentiation of B cells into plasma cells and production of IgM in the absence of external stimuli.<sup>36</sup>

Several familiarity and genetic studies have indicated that heritable components control the serum levels of IgE, IgA and IgG<sup>37–39</sup> and identified a number of loci

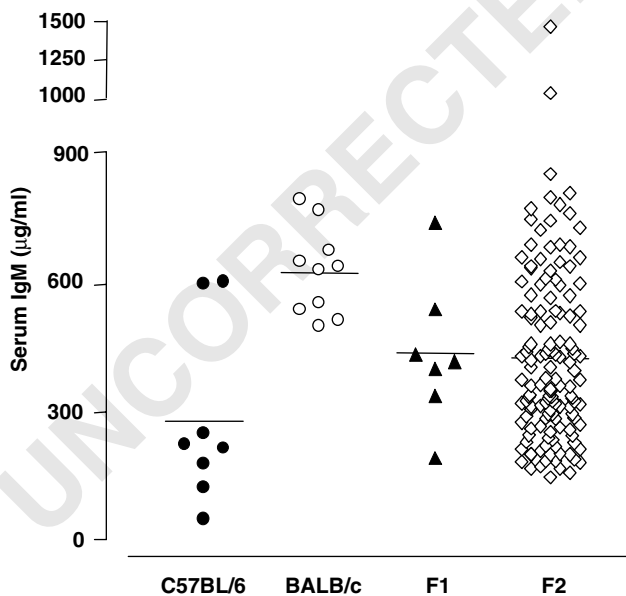
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influencing total serum levels of these antibodies in humans.<sup>40–42</sup> There is also convincing evidence for parent–offspring effects and genetic heritability in human serum IgM levels<sup>37–39</sup> as well as evidence for genetic control of natural IgM titers in mammals,<sup>43</sup> birds<sup>44</sup> and fish.<sup>45</sup> However, no linkage evidence for quantitative trait loci influencing serum IgM has been reported and the genetics of the homeostatic regulation of serum IgM remains largely unknown. Here, we characterized a total IgM serum phenotype in mouse inbred strains and performed a genome-wide screen that identified a region on mouse chromosome 13 that contained a locus (loci) controlling IgM serum levels. Searching for the cellular and genetic basis of serum IgM phenotype we correlated the level of serum IgM with the frequency of IgM-producing cells and the expression of the *Irf4* gene.

## Results

### Serum IgM phenotype

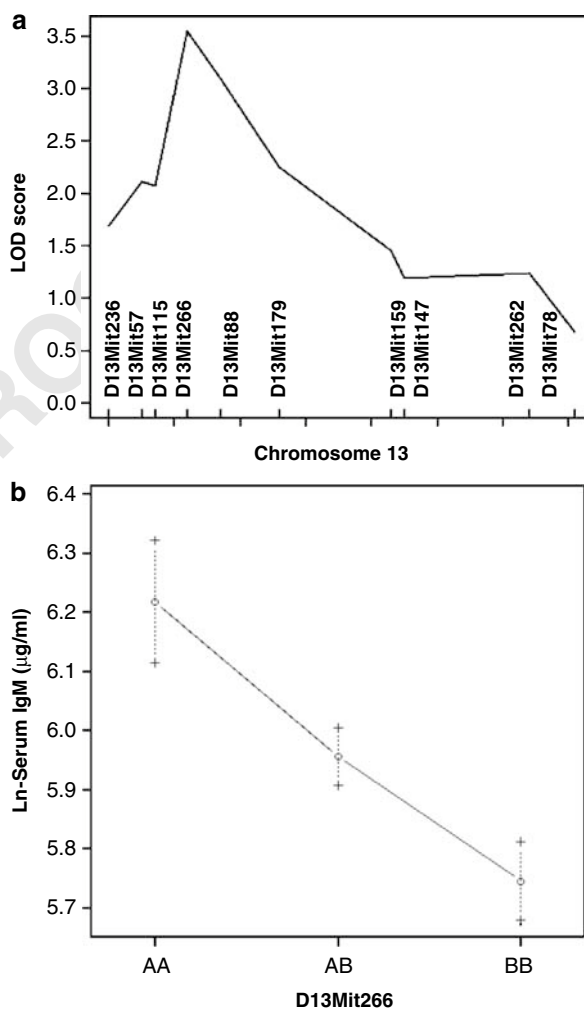
To investigate the genetic control of serum IgM we analyzed the IgM concentration in the serum of C57BL/6 and BALB/c inbred strains. We found that the serum IgM level was lower in C57BL/6 than in BALB/c mice, albeit the phenotype showed reduced penetrance (Figure 1). The F1 [C57BL/6 X BALB/c] generation showed an intermediate phenotype and an F2 [C57BL/6 X BALB/c] progeny of 136 mice showed a phenotype spectrum, from low to high values, indicating that alleles controlling the phenotype were segregating in the cross (Figure 1). The serum IgM trait, measured as the logarithm of IgM concentration, was normally distributed in the F2 [C57BL/6 X BALB/c] progeny required for quantitative trait locus (QTL) analysis (data not shown).



**Figure 1** Serum IgM levels in naive mice. IgM serum concentration was measured in eight C57BL/6 (black circles), 10 BALB/c (white circles), seven F1 [C57BL/6 X BALB/c] (black triangles) and 136 F2 [C57BL/6 X BALB/c] (white diamonds) mice by Indirect ELISA. Average concentrations are shown as horizontal bars.

### Genetic mapping of serum IgM phenotype

We performed a QTL analysis for this trait scanning the genome with 100 microsatellite markers in the 136 F2 progeny. We found that a region in proximal mouse chromosome 13 was linked to the serum IgM trait, with a 95% confidence interval between the markers D13MIT57 and D13MIT179, where the highest associated marker D13Mit266 reached a LOD score of 3.54 corresponding to an empirically estimated genome-wide *P*-value of 0.04. The chromosome 13 locus was named IgMSC1 (IgM-secreting cells locus 1; Figure 2a) and covered a region encompassing the *Irf4* gene. Although no other chromosomal region reached the level of genomic significance, suggestive LOD scores were detected in chromosome 7 at marker D7MIT318 (LOD 2.05) and in chromosome 9 at marker D9MIT53 (LOD 2.02) that could represent



**Figure 2** (a) Mapping the IgMSC1 locus. LOD score curve for mouse chromosome 13 represents the likelihood for linkage of the serum IgM trait with the represented markers. Regions of significant linkage show LOD score above 3.0. The *x* axis ticks represent the relative position of microsatellite markers along this chromosome, from left to right: D13Mit236, 4 cM; D13Mit57, 9 cM; D13Mit115, 11 cM; D13Mit266, 16 cM; D13Mit88, 21 cM; D13Mit179, 30 cM; D13Mit159, 47 cM; D13Mit147, 49 cM; D13Mit262, 68 cM; and D13Mit78, 75 cM. (b) IgM serum trait control by the IgMSC1 locus. The F2 (C57BL/6 X BALB/c) mice were classified according to their genotype at D13Mit266 and the mean trait value and the s.e. are represented for each group.

additional loci controlling the serum IgM phenotype, albeit with a smaller genetic effect as compared with IgMSC1. To evaluate the IgMSC1 genetic effect in the serum IgM phenotype F2 mice were classified according to their genotype at the highest linked marker in the IgMSC1 locus (D13Mit266). This analysis shows that the IgMSC1 locus has an additive mode of action (Figure 2b) and controls a significant fraction of the phenotypic variance (15.5%).

#### IgM-secreting cells and serum IgM

With the aim of determining the cellular basis of the IgM serum trait we started by counting the total number of B cells, sIgM<sup>+</sup> B cells and IgM-secreting cells in the spleen of C57BL/6 and BALB/c adult mice. We found no significant differences in the total number of B cells or in the total number of mature sIgM<sup>+</sup> B cells (data not shown), suggesting that regulation of B-cell spleen cellularity and B-cell maturation in the spleen is similar in these strains. However, we also found that C57BL/6 have significantly lower number of IgM-secreting cells in the spleen, offering an explanation for the observed lower serum IgM phenotype (Figure 3). Moreover, we purified B-cell populations to show that the number of IgM-secreting cells within splenic B cells was lower in C57BL/6, suggesting that these mice have less efficient plasma cell differentiation as compared with BALB/c mice (Figure 4a). These findings suggested that the cellular basis for the serum IgM trait resides in final steps of the peripheral B-cell maturation and terminal differentiation in the spleen.

It has been described that the B-cell population (B1) localizing in the pleural and peritoneal cavities is an important source of natural IgM in the mouse.<sup>46</sup> To assess the contribution of peritoneal B cells to the observed phenotype we decided to characterize the population of B cells and IgM-secreting cells in the peritoneal cavity of C57BL/6 and BALB/c mice. We found that C57BL/6 mice have a decreased number of IgM-secreting cells in

the peritoneal cavity when compared with the BALB/c strain (Figure 4b). Thus, the number of IgM-secreting B cells in the peritoneal cavity may also contribute to the cellular basis of the serum IgM trait in these mouse strains. These data indicate that the reduced number of IgM-secreting cells in the C57BL/6 mice is not specific of the spleen environment and appears to be an intrinsic genetic property of the B-cell compartment.

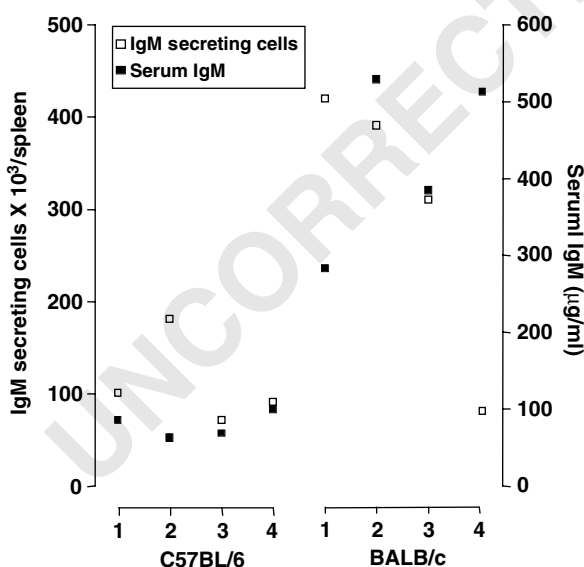
#### Irf4 gene expression

The *Irf4* gene is a key player in several stages of B-cell development, namely in controlling plasma cell differentiation<sup>30,35</sup> and co-localizes with IgMSC1 on chromosome 13. In principle, several genes in the region flanked by D13MIT115 and D13MIT88 that encompass the highest linked marker in the IgMSC1 locus could mediate the observed IgM serum phenotype. Nevertheless, the *Irf4* gene is the most plausible functional candidate in this interval as it was earlier related to antibody secretion.<sup>30,35</sup> To ascertain whether the *Irf4* gene could be involved in controlling the serum IgM trait, we quantified *Irf4* RNA by real-time PCR in purified B cells from the spleen and the peritoneal cavity. We found that *Irf4* expression was significantly reduced in splenic B cells and in peritoneal cavity B cells from the C57BL/6 (Figure 5). These data strongly suggest that *Irf4* is a candidate gene to IgMSC1 and contributes to the serum IgM phenotype by controlling the number of IgM-secreting cells.

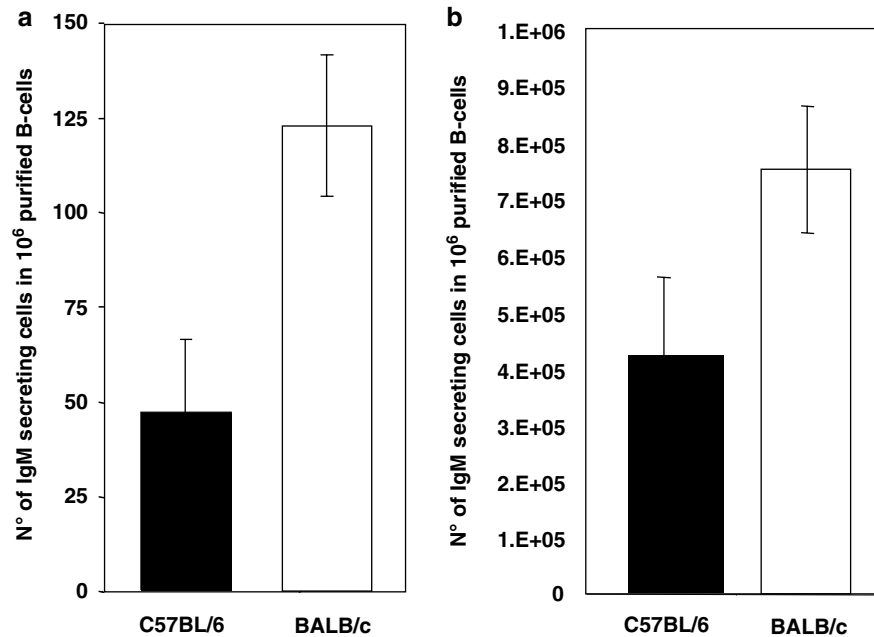
## Discussion

In this work we used a mouse genome screen approach to search for genetic factors that control the IgM levels in the serum. We identified a major locus on mouse chromosome 13 controlling the IgM serum concentration that we called IgM-secreting cells locus 1 (IgMSC1). Our results support a correlation of this phenotype with the number of IgM-producing cells in the spleen and in the peritoneal cavity, leading us to hypothesize that the IgMSC1 controls IgM serum concentration by regulating the differentiation and/or maintenance of plasma cells both in the spleen and in the peritoneal cavity. It has been described that the serum anti-RBC IgM concentration is linked to the 40–49.5 cM region on chromosome 13<sup>47</sup> and we show that gene(s) controlling natural IgM secretion are also localized in this chromosome. However, the anti-RBC locus (40–49.5 cM) and the IgMSC1 locus (9–30 cM) are somewhat apart on chromosome 13 raising the possibility that they would not correspond to the same genetic factor. We noted that the IgG serum concentration in C57BL/6 mice was significantly higher as compared with BALB/c and that IgMSC1 was not controlling levels of serum IgG in the genetic cross we analyzed (data not shown). This indicates that the genetic mechanisms that control IgM and IgG serum steady state levels have distinct and independent components.

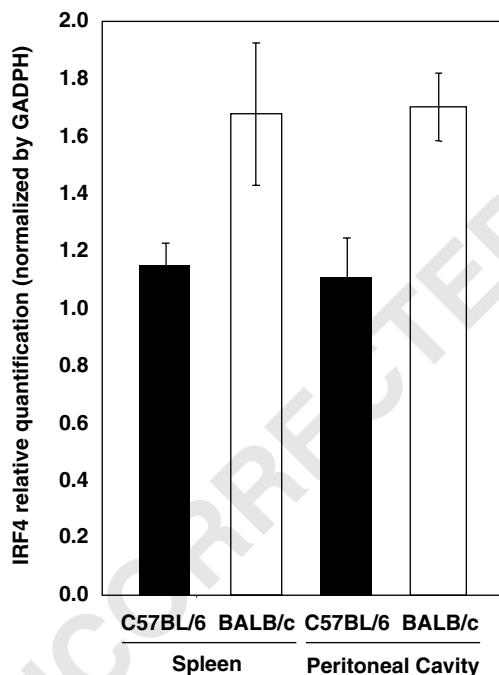
The IgMSC1 locus is tightly linked to the *Irf4* gene region on mouse chromosome 13 and we showed that the relative expression of this transcription factor is reduced in C57BL/6 B cells isolated from the spleen and from the peritoneal cavity. Thus, we propose the interferon-regulatory factor 4 as a candidate gene to mediate the IgMSC1 phenotypic effect.



**Figure 3** Number of IgM-secreting cells in the spleen correlates with serum IgM concentration. The values represent results that were obtained by ELISA-Spot-Assay and indirect ELISA in individual C57BL/6 and BALB/c mice.



**Figure 4** Decreased number of IgM secreting cell counting in purified splenic B cells and purified peritoneal cavity B cells in C57BL/6 mice. (a) Number of IgM-secreting cells in splenocyte B cells pooled from groups of eight mice. The results represent the mean and s.d. of triplicate samples assayed simultaneously. (b) Number of IgM-secreting cells in peritoneal cavity B cells pooled from groups of five mice and representing the mean and s.d. of two separate experiments assayed in triplicates.



**Figure 5** *Irf4* gene expression is decreased in purified B cells from the splenic and peritoneal cavity of C57BL/6 mice. RNA of the *Irf4* gene was quantified in purified B cells from spleens of five C57BL/6 (black bars) and five BALB/c (white bars) adult mice or in purified B cells from pooled peritoneal cavity lavages of five C57BL/6 (black column) or five BALB/c (white column) adult mice. The results represent the mean and s.d. of the average of samples assayed simultaneously.

A series of recent studies have demonstrated critical functions for IRF-4 at several stages of B-cell development, where it seems to preferentially limit clonal

expansion and promote differentiation processes.<sup>32,48</sup> *Irf4* is a member of the interferon-regulatory factor family of transcription factors characterized by a specific DNA-binding domain and has the ability to either activate or repress the transcription of interferon-inducible genes.<sup>18,31</sup> Expression of *Irf4* is restricted to cells of the immune system, including lymphocytes, dendritic cells and macrophages, in which it has been linked to a variety of functions, such as proliferation, apoptosis and differentiation.<sup>33,34,49</sup> *Irf4* knockout mice have normal early B-cell development but profound reduction in serum immunoglobulin concentrations and show incapability to mount antibody responses because of a late-stage blockage in peripheral B-cell maturation that leads to the absence of plasma cells, plasmablasts and class-switched B cells.<sup>30,35</sup> Recently, the *Irf4* gene has been demonstrated to have a biphasic expression pattern: it is expressed in immature B cells in the bone marrow,<sup>48</sup> absent from proliferating germinal center centroblasts and then re-expressed in a subpopulation of centrocytes in the germinal center and in plasma cells.<sup>50</sup> Extending these studies, the evidences provided here introduce the notion that natural genetic variation at the *Irf4* gene impacts in its expression levels, being a major factor in the quantitative control of terminal B-cell differentiation and, consequently of natural IgM serum concentration.

It is plausible that a differential effect of *Irf4* alleles could explain, at least in part, the observed differences in frequency of IgM-secreting cells and IgM serum concentration when we compared the C57BL/6 and BALB/c mouse strains. It is reported that the coding region of the *Irf4* gene in C57Bl/6 and BALB/c mice shows no sequence differences except for a synonymous point mutation in the exon 6 (NCBI entries with accession numbers AAA75309.1 and AAA75283.1). On the other hand, regulatory elements have been described in the

human *Irf4* gene encompassing the 5 kb upstream the *Irf4* coding region.<sup>51</sup> However, we have sequenced 4763 bp upstream the translational starting site of the *Irf4* gene and did not find sequence differences between C57Bl/6 and BALB/c mice. Thus, it is possible that the allelic variation controlling the observed differences in RNA expression lies in yet unidentified *Irf4* gene regulatory regions. In fact, the functional conformation of the *Irf4* promoter region suggests that its expression is strictly regulated in the chromatin context.<sup>51</sup> Detailed functional studies of the control region of the mouse *Irf4* gene would be needed to enable a systematic analysis of the differential expression of the C57Bl/6 and BALB/c *Irf4* alleles.

This work highlighted a region on mouse chromosome 13 that contains genetic factor(s) involved in the circuitry of the serum immunoglobulin homeostasis and led us to hypothesize that allelic variation in the *Irf4* gene may control the homeostasis of serum IgM in the mouse at the level of the generation of antibody-producing cells. These findings may motivate human studies on the genetic control of IgM and natural antibody levels in normal individuals and in the context of Ig deficiencies and inflammatory autoimmune diseases, particularly focused on genes mapping in the region ortholog to IgMSC1, including the *Irf4* gene.

## Materials and methods

### Mice

The mouse strains used in this study included the C57BL/6 and BALB/c as well as mice from the genetic crosses F1 (C57BL/6 X BALB/c) and F2 (C57BL/6 X BALB/c). All mice were females between 6 and 8 months of age, bred and maintained in conventional housing facilities at the Instituto Gulbenkian de Ciência. F1 (C57BL/6 X BALB/c) mice were used to generate an F2 (C57BL/6 X BALB/c) progeny of 136 animals.

### Genotyping

Genomic DNA was extracted from mouse tails with standard digestion/precipitation methods. The F2 generation was genotyped using 100 microsatellite DNA markers, polymorphic for the parental strains. Markers were chosen according to their chromosomal position as given by the Broad Institute for genome research (USA), to cover uniformly the whole mouse genome at an average density of 20 cM. The genotypes for each locus were determined by DNA marker amplification using conventional PCR techniques. Amplification products were analyzed in 4% agarose gels (Cambrex BioScience, USA) with ethidium bromide (Sigma-Aldrich, USA) and the allele size was determined using the Eagle Eye II video system (Stratagene, USA). Each individual of the F2 progeny was typed for each marker as homozygous for one or the other parental strain or as heterozygous when presenting both the parental strain alleles.

### IgM serum concentration

Standard indirect ELISA was used to quantify total IgM concentrations in mouse sera. Briefly, flat-bottomed 96-well ELISA plates were coated overnight at 4 °C, with goat anti-mouse-IgM-UNLB human adsorbed antibody (Southern Biotech., Birmingham, AL, USA) diluted in

coating buffer (0.05 M Na<sub>2</sub>CO<sub>3</sub>). The plates were washed with PBS, blocked with PBS-gelatin 1% and incubated at 37 °C for 60 min. After washing, eight serial dilutions of sera to be tested (starting at 1:90 and using 1:3 dilution steps) were incubated for 60 min at 37 °C and washed. Bound seric IgM was detected by the peroxidase-conjugated goat anti-mouse IgM-HRP human adsorbed antibody (Southern Biotech.) diluted in PBS-Gelatin 1%-Tween 0.075% (1:4000), revealed with the substrate *o*-phenylenediamine (Calbiochem, USA) and H<sub>2</sub>O<sub>2</sub> and the OD read at 490 nm. The concentrations of IgM antibody in the samples were calculated against standard curves obtained with purified monoclonal IgM (mouse-IgM UNLB, clone 11E10, Southern Biotech.) tested in the same microtiter plate.

### Genetic analysis

Quantitative trait locus analysis was performed by using the normal model in the R/QTL software.<sup>52</sup> This program calculates logarithm of odds (LOD scores) over intervals between linked markers, generating likelihood plots of genetic association with the phenotype across the genotyped chromosomal regions. The genome-wide statistical significance of the results was empirically determined by performing permutation tests<sup>53</sup> and the QTL confidence interval was calculated using the Bayesian method of R/QTL as formulated by Karl Broman in <http://www.rqtl.org/manual/html/baye-sint.html>.

### IgM-secreting cells

The number of IgM-secreting cells was determined by ELISA-Spot-Assay (ESA). The plates were coated overnight at 4 °C with goat anti-mouse IgM-human adsorbed (Southern Biotech.), diluted in coating buffer (0.05 M Na<sub>2</sub>CO<sub>3</sub>) and the plates saturated with PBS containing 1% gelatin. Serial three-fold dilutions of total splenocytes (starting with 10<sup>6</sup> cells), spleen purified B-cells (starting with 3 × 10<sup>6</sup> cells) or sorted peritoneal cavity B cells (starting with 7.5 × 10<sup>3</sup> cells) were performed in RPMI-1640 medium supplemented with 10% heat inactivated fetal calf serum (FCS), 2 mM L-glutamine, 10 mM HEPES (pH 7.4), 50 mM 2-mercaptoethanol, 100 U penicillin and 100 mg ml<sup>-1</sup> streptomycin (all from Life Technologies, NY, USA). Cell dilutions were added and the plates were sealed and incubated overnight at 37 °C in a humidified 5% CO<sub>2</sub> atmosphere. Non-adherent cells were thereafter removed by flicking the plate followed by lysis with 0.1% Tween 20 in water. After extensive washing with PBS containing 0.05% Tween 20 the plates were incubated with goat anti-mouse IgM-AP-conjugated antibody (Southern Biotech.), washed again and revealed with BCIP-AMP substrate (2.3 mM of 5-bromo-4-chloro-3-indolyl phosphate diluted in 2-amino-2-methyl-1-propanol buffer, Sigma, St Louis, MO, USA). After incubation overnight at 4 °C and washing three times with distilled water, the traces of IgM-secreting cells were revealed as blue spots and counted.

### B-cell purification

Splenocyte suspensions were prepared by straining of the spleens through a nylon mesh and peritoneal cavity cell suspensions were obtained from peritoneal lavage with 7 ml of PBS supplemented with 2% FCS. Purified B cells, with an average purity of 98%, were obtained from

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splenocyte and peritoneal cavity cell suspensions through high speed cell sorting (MoFlo, Cytomation Inc., USA), after labeling with FITC-anti-mouse CD19 (clone 1D3, BD Biosciences, CA, USA).

#### RNA isolation and real-time PCR

Total RNA from sorted splenic and peritoneal B cells was obtained using an RNeasy Mini Kit (Qiagen), following the manufacturer's protocol for animal cells. RNA was converted to cDNA (Transcriptor First Strand cDNA Synthesis Kit, Roche) using random hexamer primers. *Irf4* (Mm0051634-m1) expression was quantified using TaqMan Gene Expression Assays from ABI with TaqMan Universal PCR master mix. The gene expression quantification reactions were performed in an ABI Prism 7900HT system. Relative quantification of *Irf4* (Mm0051634-m1) in each real-time PCR was obtained after normalization for GAPDH expression measured in the same PCR and used the  $2^{-\Delta\Delta CT}$  analysis method.<sup>54</sup>

#### Statistical analysis

Statistically significant differences in serum IgM concentration, B-cell number, IgM-secreting cell number and *Irf4*-relative expression were estimated by Student's *t*-test. Differences with  $P < 0.05$  were considered significant.

## Acknowledgements

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