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Research article

**Gpnmb**<sup>R150X</sup> allele must be present in bone marrow derived cells to mediate DBA/2J glaucoma

Michael G Anderson†1,2, K Saidas Nair†3, Leslie A Amonoo1, Adrienne Mehalow3, Colleen M Trantow1, Sharmila Masli4 and Simon WM John*3,5,6

Address: 1Department of Molecular Physiology and Biophysics, University of Iowa, Iowa City, Iowa USA, 2Department of Ophthalmology and Visual Sciences, University of Iowa, Iowa City, Iowa, USA, 3The Jackson Laboratory, Bar Harbor, Maine, USA, 4Schepens Eye Research Institute and Department of Ophthalmology, Harvard Medical School, Boston, Massachusetts, USA, 5Howard Hughes Medical Institute, Bar Harbor, Maine, USA and 6Tufts University School of Medicine, Boston, Massachusetts, USA

Email: Michael G Anderson - michael-g-anderson@uiowa.edu; K Saidas Nair - si.nair@jax.org; Leslie A Amonoo - leslie-amonoo@uiowa.edu; Adrienne Mehalow - adrienne.mehalow@jax.org; Colleen M Trantow - colleen-trantow@uiowa.edu; Sharmila Masli - sharmila.masli@schepens.harvard.edu; Simon WM John* - simon.john@jax.org

* Corresponding author    †Equal contributors

Abstract

**Background:** The **Gpnmb** gene encodes a transmembrane protein whose function(s) remain largely unknown. Here, we assess if a mutant allele of **Gpnmb** confers susceptibility to glaucoma by altering immune functions. DBA/2J mice have a mutant **Gpnmb** gene and they develop a form of glaucoma preceded by a pigment dispersing iris disease and abnormalities of the immunosuppressive ocular microenvironment.

**Results:** We find that the **Gpnmb** genotype of bone-marrow derived cell lineages significantly influences the iris disease and the elevation of intraocular pressure. GPNMB localizes to multiple cell types, including pigment producing cells, bone marrow derived F4/80 positive antigen-presenting cells (APCs) of the iris and dendritic cells. We show that APCs of DBA/2J mice fail to induce antigen induced immune deviation (a form of tolerance) when treated with TGFβ2. This demonstrates that some of the immune abnormalities previously identified in DBA/2J mice result from intrinsic defects in APCs. However, the tested APC defects are not dependent on a mutant **Gpnmb** gene. Finally, we show that the **Gpnmb** mediated iris disease does not require elevated IL18 or mature B or T lymphocytes.

**Conclusion:** These results establish a role for **Gpnmb** in bone marrow derived lineages. They suggest that affects of **Gpnmb** on innate immunity influence susceptibility to glaucoma in DBA/2J mice.

Background

The glaucomas are a common group of potentially blinding diseases that by 2010 will affect approximately 60 million people worldwide [1]. The glaucomas share a clinical phenotype including a progressive degeneration of the optic nerve [2,3]. This glaucomatous optic neuropathy
causes a progressive and irreversible loss of vision, and may lead to complete blindness. Significant known risk factors for glaucoma include elevated intraocular pressure (IOP), aging, positive family history, race, abnormal optic nerve head morphology and decreased central corneal thickness [4-12]. Of these, the only currently modifiable risk factor is IOP, which is the target of all existing glaucoma treatments [13-15]. One means of gaining a better understanding of glaucoma pathogenesis, and ultimately the creation of new therapeutic interventions, is to study the underlying molecular pathways with experimental systems such as the mouse. With the recent descriptions of mouse strains and techniques relevant to studying glaucoma, genetic approaches in mice offer great promise for testing new and potentially novel hypotheses related to glaucoma [16-19].

Experiments with DBA/2 (D2) mice have suggested that immune abnormalities may contribute to some forms of glaucoma [20,21]. D2 mice develop a form of glaucoma involving a pigment dispersing iris disease that aberrantly deposits pigment throughout the anterior chamber, including the drainage structures of the eye [22-24]. As a consequence, D2 and several closely related strains develop elevated IOP and glaucomatous neuropathy [24-30]. Eyes of D2 mice also exhibit multiple abnormalities in ocular immune privilege [20], including deficient anterior chamber associated immune deviation (ACAID). Importantly, the iris pigment dispersion component of the D2 iris disease and the inability to support ACAID are simultaneously rescued when their marrow is repopulated with cells from B6D2F1 mice [20]. While the bone marrow origin of immune cells involved in ACAID explains the restoration of ACAID by B6D2F1 bone marrow cells (BMC), the simultaneous resolution of the pigment dispersing iris disease links this disease to bone marrow derived cells. The B6D2F1 mice that served as a source of normal BMC are offspring of a cross between glaucoma prone D2 and normal C57BL/6J (B6) mice. These B6D2F1 mice are heterozygous for all B6 and D2 specific alleles across the entirety of the autosomal genome. Therefore, no specific genes were mechanistically implicated in the recovery of ACAID or the iris disease. The goal of the current experiments is to identify the gene(s) responsible for mediating this BMC contribution to D2 phenotype.

Genetic experiments have previously shown that mutations in two genes digenically promote glaucoma in D2 mice, Tyrp1 and Gpnmb [22,23]. The Tyrp1 gene encodes a relatively well characterized melanosomal enzyme that is required for eumelanogenesis and is localized at the melanosomal membrane. So far there are no reports of either expression or influence of this gene on cell types derived from the bone marrow. Therefore, Tyrp1 is unlikely to be relevant in the BMC mediated glaucoma phenotype of D2 mice. Less is known concerning Gpnmb. The Gpnmb gene is predicted to encode a transmembrane protein with homology to the melanosomal protein, silver (pMel17), but the function(s) of GPNMB remain largely unknown. Interestingly, Gpnmb is expressed by some BM derived cell types [31]. Recent studies demonstrate that GPNMB is expressed by macrophages and functions as a negative regulator of macrophage mediated inflammatory responses [32]. In another study, it was reported that GPNMB binds activated T cells. This binding causes inhibition of TCR-induced T-cell activation for both primary and secondary immune responses [33]. Thus, Gpnmb is an attractive candidate that potentially influences the BMC mediated phenotypes in D2 mice.

Here, we test the hypothesis that the Gpnmb gene contributes to the bone marrow dependent events in D2 glaucoma. Using D2 mice that only differ in Gpnmb genotype [34], we demonstrate that BMC lineages containing wild-type Gpnmb can rescue both the pigment dispersing iris disease and IOP elevation of D2 mice. Because GPNMB protein localizes to BM derived antigen-presenting cells (APCs) of the iris and to cultured dendritic cells, we assessed potential phenotypes that may be modulated by these APCs. First, we find that APCs of D2 mice are abnormal in that they fail to induce ACAID, this effect is not dependent on Gpnmb. Second, we examined aqueous humor levels of IL18. Bone marrow derived APCs in the iris are a potential source of IL18 [35] and IL18 is reported to become elevated in the aqueous humor of D2 mice as disease progresses [21]. In contrast to the previous report [21], we find no evidence for involvement of IL18 in the D2 disease. Surprisingly, we finally show that effectors of adaptive immune responses, mature B or T lymphocytes, are not necessary to mediate the glaucoma inducing iris disease of D2 mice.

Results

Gpnmb is a candidate for mediating BM derived contributions to D2 ocular disease

We have previously shown that the D2 iris disease is strongly influenced by a BMC lineage [20]. Our consideration of candidates capable of mediating this influence was based in part on previous genetic experiments demonstrating that the D2 iris disease is initiated by the digenic interaction of the Tyrp1 and Gpnmb genes [22,23]. To ascertain whether either Tyrp1 or Gpnmb are plausible candidates for mediating the BMC contributions to D2 iris disease, semi-quantitative RT-PCR was performed to compare gene expression patterns of these genes in iris and bone marrow. Since valid candidates would likely be expressed in cells of the immune system, their expression levels in lymph nodes and thymus were also tested. Supporting a potential role for Gpnmb in bone marrow derived phenotypes, expression of Gpnmb was detected in
all of these tissues. In contrast, the expression of *Tyrp1* was limited to only the iris (Fig. 1).

Having found that *Gpnmb* was expressed in the appropriate tissues, we next assessed the nature of the *Gpnmb*<sup>R150X</sup> mutation. The *Gpnmb*<sup>R150X</sup> mutation in D2 mice creates a premature stop codon. As a result of nonsense-mediated mRNA decay, the mutation is predicted to result in a severe decrease in *Gpnmb* transcript levels [36]. To test this, a quantitative real-time PCR assay for *Gpnmb* was performed. Expression in irides of young predisease D2 mice homozygous for the *Gpnmb*<sup>R150X</sup> mutation was compared to age and sex matched wild-type controls. With samples normalized to levels of *Rn18s*, the *Gpnmb*<sup>R150X</sup> mutation resulted in a severe reduction in *Gpnmb* transcript levels (~18 fold, data not shown). In agreement with this, no protein with the expected molecular size was detected by Western analysis using a GPNMB antibody (Fig. 1B). Similar analysis of *Tyrp1* levels demonstrated no significant change in transcript levels. The expression of *Gpnmb* in immune tissues and the absence of GPNMB protein in R150X mutants support the hypothesis that this mutation disrupts BMC lineage dependent functions of GPNMB and that it may be responsible for bone marrow mediated aspects of the D2 disease.

**A functional requirement for Gpnmb in bone marrow**

To genetically test whether the GPNMB genotype of BM derived lineages influences the D2 iris disease, we took advantage of the DBA/2J substrain, D2-*Gpnmb<sup>+</sup>* that has a wild-type *Gpnmb* allele but no other known differences to modern D2 mice [34]. Whereas D2 mice with the *Gpnmb*<sup>R150X</sup> mutation develop an iris disease characterized by significant pigment dispersion, iris atrophy, and anterior chamber enlargement, irides of D2-*Gpnmb<sup>+</sup>* mice exhibit far milder phenotypes (Fig. 2A–J). Using these mice as donors and recipients in bone marrow transfers, we next asked whether the genotype of *Gpnmb* in BMC lineages was capable of influencing iris phenotypes. Lethally irradiated D2 mice were reconstituted with D2-*Gpnmb<sup>+</sup>* bone marrow and followed clinically for indications of iris disease (Fig. 2K–O). In all figures and text, the genotype of donor bone marrow is written first followed by an arrow and then the recipient genotype. For example, WT → R150X where D2-*Gpnmb<sup>+</sup>* (WT) bone marrow was transferred into D2 mice homozygous for the R150X mutation. At ages when the D2 iris disease is normally severe, iris phenotypes in D2 mice reconstituted with D2-*Gpnmb<sup>+</sup>* bone marrow were significantly rescued toward the wild-type iris phenotype. They developed less pigment dispersion, less transillumination, and less change in the dimensions of the anterior chamber as compared to both unmanipulated D2 mice (compare Fig 2K–O with 2F–J) and D2 mice that were reconstituted with standard D2 marrow that had the *Gpnmb*<sup>R150X</sup> mutation [20,37]. The converse experiment reconstituting lethally irradiated D2-*Gpnmb<sup>+</sup>* mice with standard D2 bone marrow (R150X → WT) was also performed (Fig. 2P–T). Iris phenotypes of D2-*Gpnmb<sup>+</sup>* mice reconstituted with standard D2 bone marrow were unaltered and maintained an iris indistinguishable from unmanipulated D2-*Gpnmb<sup>+</sup>* mice (compare Fig. 2P–T with 2A–E). In sum, these results indicate that the *Gpnmb*<sup>R150X</sup> mutation results in the bone-marrow derived portion of the D2 iris disease. However, the influence of the *Gpnmb*<sup>R150X</sup> mutation acting via bone marrow derived lineages is not itself sufficient to induce the iris disease in otherwise healthy mice.
IOP elevation in D2 mice can be modulated by Gpnmb bone marrow genotype

In unmanipulated D2 mice, the depigmenting iris disease is typically followed by IOP elevation and enlargement of the anterior chamber. Because the iris disease of D2 mice reconstituted with D2-Gpnmb⁺ bone marrow is amelio-

Figure 2
D2-Gpnmb⁺ bone marrow suppresses iris depigmentation and anterior chamber enlargement. Representative eyes of the indicated strains and ages are shown. The three left most columns show broad beam illumination to assay for the presence of dispersed pigment within the anterior chamber and iris stromal morphology. The fourth column shows a transilluminating view assaying the degree of iris depigmentation, detectable as red areas within the image where reflected light is passing through the iris. The fifth column show anterior chamber dimensions. Each row represents a different genetic context of either unmanipulated mice (rows 1 & 2) or bone marrow chimeras (rows 3 & 4). (A to E) Unmanipulated D2-Gpnmb⁺ mice (WT) exhibit a characteristic iris stromal atrophy phenotype caused by the Tyrp1b mutation that is largely devoid of significant pigment dispersion at all examined ages (1–15 mo). Eyes of D2-Gpnmb⁺ mice do not develop significant transillumination and maintain a normal anterior chamber depth with very little space between the iris and cornea. (F to J) Unmanipulated D2 mice (homozygous for the GpnmbR150X mutation) develop a severe pigment dispersing iris disease. At 5–6 mos, the initial stages of iris disease are apparent by a slight peripapillary thickening and subtle changes to the morphology of the iris stroma (note slightly roughened appearance). Subsequently, dispersed pigment becomes aberrantly deposited on a variety of structures including the surface of the iris, lens, and cornea. Dispersed pigment is also deposited in the aqueous humor drainage structures, leading to increased IOP and enlargement of the anterior chamber (space indicated by arrows in J). (K to O) D2-Gpnmb⁺ marrow transferred to D2.GpnmbR150X recipients (WT → R150X) exhibit a pronounced suppression of the typical D2 disease progression. Most notably, the extent of iris degeneration is lessened (compare the peripapillary region of M vs H, and degree of transillumination defects in N vs I) and the anterior chamber does not become enlarged (compare O vs J). (P to T) D2 marrow transferred to D2-Gpnmb⁺ recipients (R150X → WT) exhibit no alterations in ocular phenotype compared to unmanipulated D2-Gpnmb⁺ mice.
Bone marrow genotype affects glaucomatous IOP elevation. To directly test this, IOP was measured at multiple ages for bone marrow chimera cohorts and unmanipulated D2 mice of differing Gpnbm genotype (Fig. 3). 10-mo D2-Gpnbm+ mice had a relatively normal mean IOP of 16 mmHg, 10-mo unmanipulated D2 mice had an elevated mean IOP of 20 mmHg, and 10-mo D2 mice reconstituted with D2 marrow (R150X → R150X – a syngeneic control for possible influence of bone marrow transfer itself) also had an elevated IOP of 20 mmHg.

Importantly, 10-mo D2 mice reconstituted with D2-Gpnbm+ bone marrow (WT → R150X) had a mean IOP of 16 mmHg, indicating that the influence of Gpnbm in bone marrow was able to prevent IOP elevation from occurring. Furthermore, the IOP of D2 mice reconstituted with D2-Gpnbm+ bone marrow (WT → R150X) remained at these normal levels to at least 16-mo. These results further support the conclusion that the GpnbmR150X mutation influences the glaucoma phenotype of D2 mice via BMC lineages.

Figure 3
Bone marrow genotype affects glaucomatous IOP elevation. Plot of IOP vs Gpnbm genotype, with age and numbers of mice indicated. D2-Gpnbm+ mice exhibit an average IOP typical of many non-glaucomatous mouse strains. As previously described, D2 mice (homozygous for the GpnbmR150X mutation) exhibit an elevated IOP at 10 mo of age, as do D2 mice receiving syngeneic bone marrow transfers (R150X → R150X). Chimeric mice in which D2-Gpnbm+ bone marrow has been transferred into D2 hosts (WT → R150X) continue to maintain non-glaucomatous IOP values, whether examined at 10 mo, or beyond. The IOPs of both Gpnbm+ mice and chimeric mice with Gpnbm+ bone marrow were significantly lower than those of all mice with GpnbmR150X mutant marrow (P < 0.002 for all comparisons at various ages, t test).

GPNMB localization within F4/80 positive APCs

Relatively little is known concerning the Gpnbm gene, or its orthologs in other species [38-40]. Although the ocular localization of GPNMB protein has not previously been described, experiments have suggested two ocular cell types in which GPNMB would likely be found. First, GPNMB is likely to be present in melanin producing cells [22,41,42]. Second, GPNMB is likely to be present in BM derived APCs, including macrophages and a subset of dendritic cells (DCs) [31,32]. In order to begin identifying specific cell types in which Gpnbm may function, an anti-GPNMB antibody was used to localize GPNMB in normal adult C57BL/6J mouse eyes (Fig. 4). In adult C57BL/6J mice, we found that GPNMB is robustly detectable within multiple pigmented cells of the eye, including: melanocytes of the iris stroma and pigmented cells of the iris pigment epithelium, the pigmented epithelia of the ciliary body (Fig. 4 A-D), the choroid and to a lesser extent the retinal pigment epithelium (Fig. 4E, F). Diffuse low level labeling was observed throughout the neural retina. GPNMB was not detected in other ocular tissues such as the cornea or lens. In each of the pigmented tissues, GPNMB localization was punctate, likely as a consequence of localization within melanosomes. Though important for other aspects of Gpnbm function [43], there is no evidence that any of the pigment producing cells of the eye are derived from the bone marrow. Thus, we next examined whether GPNMB is present in ocular APCs.

The iris normally contains a robust population of APCs [44] and GPNMB has previously been demonstrated to be present in mouse DCs [31]. To determine if a portion of GPNMB in the iris is within APCs, the APC marker F4/80 was tested for co-localization with GPNMB protein. In adult C57BL/6J mice, GPNMB was indeed present within the cytoplasm of iridial F4/80 positive cells (Fig. 4G). To assess the localization of GPNMB in DC lineages, bone marrow progenitors were grown in cell culture, stimulated to differentiate into DCs, and utilized for immunohistochemistry. In DC cells grown in these conditions, GPNMB was observed in intracellular granules (Fig. 4H). Because F4/80 positive APCs and DCs are BM derived cell lineages, this result indicates that the bone marrow functions of Gpnbm are likely to be mediated, in part or in whole, by these cells.

Gpnbm deficient APCs fail to induce immune deviation

We next determined the effect of the Gpnbm mutation on the ability of APCs to induce immune deviation. The healthy eye exists in a state of immunologic balance that provides immune protection by decreasing risk of immunopathogenic injury to ocular tissues [45]. Anterior chamber associated immune deviation (ACAID) is an anti-inflammatory mechanism that is a form of eye-dependent tolerance mediated by ocular APCs. The immune devia-
tion is initiated by F4/80+ ocular APCs that capture antigen in the eye and migrate via the blood to the spleen [46]. At that site, these APCs generate a population of regulatory T cells capable of inhibiting a Th1-mediated inflammatory immune responses such as a delayed type hypersensitivity response (DTH). We have previously demonstrated that D2 mice lack the ability to support ACAID as evidenced by their inability to inhibit a DTH response, but the genes responsible for this finding are not defined [20].

Non-ocular APCs, such as F4/80+ macrophages derived from thioglycollate-elicited peritoneal exudate cells (PECs), can be converted into APCs that induce immune deviation similar to that induced by ocular APCs by exposing them to TGFβ2 in culture [47-50]. Thus, TGFβ2-exposed peripheral APCs resemble ocular APCs in their functional phenotype. To test the Gpnmb dependency of this APC phenotype, we compared the ability of APCs with differing Gpnmb genotypes to induce immune deviation as detected by inhibition of DTH. To allow experiments with APCs in genetically identical hosts, all experiments used B6D2F1 mice as recipients. TGFβ2-treated APCs from control B6 mice with a wild-type Gpnmb allele successfully induced immune deviation that led to inhibition of DTH (Fig. 5A). In contrast, APCs from both Gpnmb deficient and sufficient D2 mice failed to induce immune deviation when treated with TGFβ2 (Fig. 5A). Thus, the inability of D2 APCs to mediate immune deviation is not dependent on Gpnmb alone. This suggests that some property of D2 APCS other than the GpnmbR150X mutation is responsible for their inability to induce immune deviation.

In parallel with these experiments, we tested the ability of TGFβ-treated APCs from B6 Tyrp1CpGpnmbR150X congenic mice to induce tolerance. These congenic mice have the D2-derived Tyrp1CpGpnmbR150X mutations on a B6 genetic background (Methods). Also, we evaluated the expression of a panel of genes for which coordinate changes in expression normally contribute to the immune deviation inducing phenotype of APCs [51-53]. Similar to D2 APCs, the B6.Tyrp1CpGpnmbR150X derived APCs failed to induce immune deviation when treated with TGFβ2 (Fig. 5A). Arguing against a causative role for the Tyrp1 mutation, BALB/c mice have the same Tyrp1 mutation but do not have deficiencies in ACAID [20]. Together, our results suggest that a gene(s) that resides in one of the D2-derived chromosomal regions may be important in mediating APC induced tolerance. (D2-derived chromosomal regions surround the Tyrp1 and Gpnmb genes in the B6 congenic mice [43]). In agreement with this, the coordinated pattern of changes in gene expression that support immune deviating properties of APCs exhibited a classic tolerizing pattern in B6 mice, but not in D2 (Figure 5B).

**Figure 4**

GPNMB localizes to multiple pigmented tissues of the eye and ocular APCs. Cryosections of whole eyes from wild-type C57BL/6J mice were imaged in brightfield or phase illumination. GPNMB protein localization was imaged by fluorescent immunohistochemistry and confocal microscopy (red). GPNMB is present in: (A, B) multiple pigmented tissues of the eye, (C, D) iris and pigmented epithelia of the ciliary body, and (E, F) choroid. (G) Iris of normal C57BL/6J mouse showing GPNMB (red) within cells labeled by F4/80 (green). An 8 micron Z-series composite demonstrating F4/80 labeling the border of a single cell (dashed white circle) containing GPNMB. (H) BM derived dendritic cell from D2-Gpnmb+ marrow differentiated in cell culture showing punctate expression of GPNMB. (I, J) Negative controls were labelled and imaged with identical conditions and exhibited negligible signal. (I) Eyes from D2 mice (homozygous for GpnmbR150X) have no signal for GPNMB, an inset on right hand corner shows the brightfield image of the same section. (J) BM derived dendritic cells from D2 mice have no staining for GPNMB. Scale bars A-F, I = 500 μm, G, H, J = 10 μm.
Gpnmb genotype influences IL18. Secretion of IL18 by macrophages has been implicated in both Th1 and Th2 immune responses. Aqueous humor levels of the immune modulating cytokine IL18 are reported to become elevated in D2 mice, and the levels of IL18 were suggested to associate with increases in IOP [21]. To test whether Gpnmb genotype influences IL18 levels, aqueous humor was collected from cohorts of either D2 or B6.Typr1bGpnmbR150X mice homozygous for the GpnmbR150X mutation and compared to the strain matched controls, D2-Gpnmb+ or C57BL/6J (Fig. 6A). Additionally, expression of IL18 mRNA in tissue enriched in ciliary body was analyzed by real-time PCR (Fig. 6B). In these mice, there was no correlation between Gpnmb genotype and IL18 levels. Furthermore, in contrast to previous studies of IL18 in D2 mice [21], we also found no correlation between IL18 and age. Indeed, in our studies, IL18 levels were significantly lower in 9 mo old D2 mice compared to 3 and 6 mo old D2 mice ($P < 0.05$). These results fail to support a role for IL18 in the D2 form of glaucoma and suggest that the relevant bone marrow derived functions of Gpnmb do not influence this aspect of ocular immunity.

**Deficiency of Rag1 does not affect Gpnmb mediated disease**

GPNMB is present within APCs and APCs initiate and direct adaptive immune responses. Therefore, we tested whether mature B or T lymphocytes, play a role in the Gpnmb mediated iris disease. To accomplish this, we exploited a mutation in the recombination activating gene Rag1. Mice deficient for Rag1 are characterized by arrested development of T and B cells and are therefore incapable of adaptive immune reactions mediated by mature T or B cells. To test the necessity of T and B cells for the pigment dispersing iris disease, we utilized B6.Typr1bGpnmbR150X mice [43] with a Rag1 mutation. As a consequence of the D2-derived Typr1b and GpnmbR150X mutations, B6.Typr1bGpnmbR150X mice develop an iris disease with near identical features to that of D2 mice [43]. Ablating the adaptive immune functions of T and B cell had no effect on the iris disease of these mice. The iris disease of Rag1 deficient B6.Typr1bGpnmbR150X mice is indistinguishable to that of their littermates with a functional Rag1 gene and an intact adaptive immune system (Fig. 7). Similar results were obtained with mice homozygous for the Prkdcscid mutation that also lack mature B and T lymphocytes [see Additional File 1]. These experiments demonstrate that the loss of adaptive immune reactions has no influence on the age of onset, rate of progression, or final severity of the iris disease mediated by Typr1 and Gpnmb mutations. Given the dependence of disease phenotypes on the Gpnmb genotype of bone marrow but not on adaptive immunity, these results suggest that GPNMB may impact innate immune processes. Therefore, alterations of
innate immunity may participate in the pathogenesis of the iris disease.

**Discussion**

D2 mice develop a pigmentary form of glaucoma involving a pigment dispersing iris disease, increased IOP, and degeneration of retinal ganglion cells [22-24,26]. In recent years, we have begun to study early events within the anterior chamber that initiate the iris disease and IOP elevation [20,22,23,43] as well as later events associated with the neurodegeneration [37,54,55]. We have previously identified two genes responsible for the initiation of this disease, Tyrp1 and Gpnmb, both of which encode melanosomal proteins [22,42,56]. We have also shown that a non-melanosomal component contributes to the D2 iris disease which acts via a BMC lineage [20]. Here, we have identified the Gpnmb gene as an important gene influencing these bone marrow derived contributions.

![Figure 6](image-url)

**Figure 6**

**IL18 levels do not become elevated in D2 aqueous humor or ciliary body.** (A) IL18 levels in aqueous humor of age matched D2, D2-Gpnmb<sup>+</sup>, B6.Tyrp1<sup>+</sup>Gpnmb<sup>150X</sup>, and C57BL/6j mice were determined by ELISA. IL18 levels are expressed as mean +/- SEM from 5 aqueous humor samples in each group. (B) Quantitative RT-PCR analysis comparing expression levels of IL18 transcript in ciliary body enriched dissections of 10.5 mo old D2, D2.Gpnmb<sup>+</sup>, and B6.Tyrp1<sup>+</sup>Gpnmb<sup>150X</sup> mice. The threshold cycle was calculated with normalization to Rn18s and values are expressed as mean +/- SEM, with 3–5 samples in each group.

![Figure 7](image-url)

**Figure 7**

**Loss of adaptive immune reactions has no influence on Gpnmb mediated iris disease.** The Rag1<sup>tm1Mom</sup> mutation acts recessively to result in a loss of adaptive immune responses due to lack of mature B and T lymphocytes. Stocks carrying this mutation were bred to B6.Tyrp1<sup>+</sup>Gpnmb<sup>150X</sup> mice. Triple mutant mice were isolated and aged, with typical eyes of indicated genotypes shown. The analysis strikingly demonstrates that the immune deficiency associated with Rag1 mutation has no effect on Gpnmb mediated iris disease. (A) Eyes of young B6 mice with wild-type Tyrp1, Gpnmb, and Rag1 alleles have healthy irides. (B) At 15 mo, mice mutant for Tyrp1 and Gpnmb exhibit iris disease characterized by significant pigment dispersion and iris atrophy. n = 10 eyes, age range 14–16 mo. (C) At 15 mo, immune deficient mice mutant for Tyrp1, Gpnmb, and Rag1 exhibit an iris disease indistinguishable from immune competent mice on the same genetic background. N = 26 eyes.
From the sum of prior and present experiments (see Additional File 2 and refs [20,22]), an attractive hypothesis suggests that the D2 iris disease initially involves a melanosomal defect mediated by the Tyrp1 and Gpnmb genes that mildly damages the iris and causes cellular debris, including pigment, to be shed into the anterior chamber. Due to deficiency of GPNMB, a bone marrow derived lineage(s) that would normally express Gpnmb then reacts abnormally to the iris debris. This results in an inflammatory attack on the iris, severe iris atrophy and the ensuing glaucoma. The abnormally responding bone marrow derived lineage may either fail to inhibit ocular inflammation or may actively promote an inflammatory attack on the iris. This model accounts for the ability of B6D2F1 [20] or D2-Gpnmb+ (Fig. 2) bone marrow to suppress the D2 iris disease (because bone derived cells with wild-type Gpnmb alleles do not respond abnormally to the debris resulting from the melanosomal insults and mild iris damage). Additionally, this model explains why bone marrow from D2 mice with the Gpnmb<sup>R150X</sup> mutation is not sufficient to confer the severe iris disease when transplanted into recipient mice with wild-type Gpnmb alleles (because the recipients with wild-type Gpnmb alleles do not have sufficient iris damage to prime the immune system attack of the iris.). Thus, this model, explains why the bone marrow derived function of Gpnmb is a necessary, but not sufficient, component of the overall disease.

The functions of Gpnmb in BMC derived lineage(s) that are relevant to DBA/2J glaucoma remain unknown. Although Gpnmb genotype did not influence the ability of D2 APCs to mediate tolerance to OVA when they were first exposed to TGFβ2 and OVA in culture, GPNMB may still modulate some aspects of APCs that contribute to D2 phenotype and were not assessed. These functions may include their ability to process and present antigens derived from damaged iris cells, their ability to secrete various chemokines that recruit proinflammatory cells or their ability to respond to anti-inflammatory molecules in the ocular microenvironment. Regardless, we show that the immune abnormalities contributing to the Gpnmb mediated iris disease do not require functions of the adaptive immune system. This suggests that GPNMB deficiency may influence innate immune cells that are of bone marrow origin and are likely to respond to abnormalities in the iris. Experimental expression of GPNMB in activated macrophage causes reduced release of proinflammatory cytokines such as IL6, IL12 and TNF alpha [32]. We localize GPNMB within APCs. Therefore, it is possible that the Gpnmb<sup>R150X</sup> mutation alters APCs so that they are more prone to mediate inflammation and this abnormal phenotype promotes ocular inflammation in D2 mice.

Other groups have previously localized GPNMB in macrophages and in the MHC class II compartment of DCs [31].

The punctate localization of GPNMB in pigmented cells and its melanosomal targeting sequence are consistent with another function of GPNMB in melanosomes [22]. Interestingly, melanosomes and the MHC class II compartment are both lysome-related organelles [57,58]. This correlation suggests that similar to molecules such as BLOC complexes and LYST [59,60], GPNMB may be a molecule that functions in multiple classes of lysosome-related organelles.

The glaucomatous potential of the DBA/2J iris disease has recently been shown to be strikingly sensitive to genetic background [43]. When transferred to the C57BL/6J genetic background, the Gpnmb<sup>R150X</sup> and Tyrp1<sup>b</sup> mutations result in a severe pigment dispersing iris disease that is phenotypically indistinguishable from the iris disease of D2 mice. Surprisingly, however, these B6 mice are resistant to IOP elevation and glaucoma. Thus, there are additional modifier genes whose alleles determine whether or not IOP becomes elevated and glaucoma ensues. Although these modifier genes remain to be identified, our preliminary data suggest existence of at least 3 genetic loci, on different chromosomes to Gpnmb and Tyrp1, that impact the nature of the anterior chamber disease and determine if IOP becomes elevated. Given the dependence of the iris disease on bone marrow genotype, at least some of these modifiers are likely to influence immune functions. Since the chromosomal region containing either Gpnmb or Tyrp1 transferred the inability of APCs to mediate tolerance to the B6 congenic strain, there must be another strain specific allele that influences APC functions. Efforts to identify these strain specific modifier alleles are underway and are likely to shed additional light on the molecular basis of immune contributions to this glaucoma. Characterizing these genes is expected to improve understanding of APC responses within the anterior chamber and mechanisms of inducing tolerance.

**Conclusion**

We have demonstrated that Gpnmb influences the glaucoma phenotype of D2 mice by a BM derived mechanism that does not require adaptive immunity. Although immunity has not yet been shown to contribute to human pigmented glaucoma, the inflammation is subclinical in the mice and may have been undetectable on standard examination of patient’s eyes. Thus, similar mechanisms of mild inflammation may be active in human pigmented glaucoma and perhaps other ocular diseases in which pigment dispersal occurs within the anterior chamber. These findings represent an important step toward gaining a molecular understanding of the mechanisms active in this pigmented form of glaucoma.
D2 mice were obtained from The Jackson Laboratory, Bar Harbor, Maine. Some studies also utilized B6.D2-Tyrp1\(^Gpnmb^{R150X/Sj}\) mice (referred to as B6 Tyrp1\(^Gpnmb^{R150X/Sj}\)), an N10 congenic strain of mice which are homozygous for chromosomal intervals containing the D2-derived Tyrp1\(^Gpnmb^{R150X/Sj}\) alleles that have been backcrossed into the C57BL/6J genetic background [43]. Mice with a recombination activating gene 1 mutation \((Rag^{1m1Mom})\) were obtained from The Jackson Laboratory and crossed to the B6.Tyrp1\(^Gpnmb^{R150X/Sj}\) strain to produce mutants that were homozygous for all three mutations \((B6.Cg-Rag^{1m1Mom}/Tyrp1^{Gpnmb^{R150X/Sj}},\) referred to as \(Rag1\) mutant B6 Tyrp1\(^Gpnmb^{R150X/Sj}\)). Two strains of control mice with wild-type \(Gpnmb\) alleles were utilized in these studies, DBA/2J with a wild-type \(Gpnmb\) allele (DBA/2J-Gpnmb\(^Sj\), referred to as D2-Gpnmb\(^Sj\)) [34] and C57BL/6J. Since the \(Gpnmb\) allele is the ancestral D2 allele, the \(Gpnmb^{R150X/Sj}\) mutation is the only known genetic difference between the D2 and D2-Gpnmb\(^Sj\) strains [22]. C57BL/6J mice (with wild-type \(Tyrp1\) and \(Gpnmb\) alleles) were obtained from The Jackson Laboratory. All D2 background mice were maintained on a 6% fat NIH 31 diet provided \(ad\) \(libitum\), and the water was acidified to pH 2.8–3.2. All B6 background mice were similarly maintained but on an NIH 31 diet with 4% fat content. Mice were housed in cages containing white pine bedding and covered with polyester filters. The environment was kept at 21°C with a 14-h light:10-h dark cycle. All animals were treated according to the guidelines of the Association for Research in Vision and Ophthalmology for use of animals in research. All experimental protocols were approved by the Animal Care and Use Committee of The Jackson Laboratory and cross-referred to the University of Iowa.

**Generation and analysis of bone marrow chimeras**

Bone marrow chimeras were generated as follows: 4–8 wk old female D2 and D2-Gpnmb\(^Sj\) mice were lethally irradiated (1000 rad from a \(^{137}\)Cs source) and then received 200 μl intravenous injections containing \(5 \times 10^6\) T-cell depleted bone marrow cells from the indicated donor strains. Donor marrow was depleted of T lymphocytes with 10 μg/mL of purified monoclonal antibodies to CD4 (GK1.5; The Jackson Laboratory Flow Cytometry Service, Bar Harbor, Maine) and CD8a (53–6.72; The Jackson Laboratory Flow Cytometry Service). Eyes of chimeras were clinically assayed at 2–3 mo intervals.

**Intraocular pressure measurement**

IOP was measured using the microneedle method as previously described in detail [61,62]. Because the IOPs of B6 mice are very consistent, B6 mice were interspersed with experimental mice during all experiments as a methodological control to ensure proper equipment calibration and performance.

**Slit-lamp examination**

Eyes were examined with a slit-lamp bio-microscope and photographed with a 40× objective lens. Phenotypic assessment of iris disease was determined by indices of iris atrophy dispersed pigment and transillumination defects following previously described criteria [20,22-24]. Transillumination is an assay of iris disease whereby reflected light passing through depigmented areas of iris tissue are visualized as red light.

**RNA Isolation and Quantitative RT-PCR**

Tissues were dissected, homogenized, and total RNA extracted. cDNA was generated using 300 ng of total RNA isolated from dissected irides or 70 ng of total RNA isolated from ciliary body enriched dissections. Real-time PCR data were collected utilizing standard reaction conditions, with primer efficiencies determined from serial dilutions of cDNA and relative expression calculated as previously published [63-65]. Reaction conditions and primer sequences are available upon request.

**Western blotting**

Iris tissue from D2 and D2-Gpnmb\(^Sj\) was homogenized in lysis buffer (10 mM Tris pH 7.6, 150 mM NaCl, 1% Triton X-100 and supplemented with protease inhibitors). The lysate was resolved on a 4–20% SDS-PAGE gradient gel. Proteins were blotted to a PVDF membrane, followed by incubation with anti-GPNMB antibody (R&D systems), secondary antibody and detected using SuperSignal West Pico chemiluminescent substrate (Pierce Biotechnology).

**Immunohistochemistry**

Enucleated eyes were embedded in Optimal Cutting Temperature embedding medium (Tissue-Tek O.C.T. Compound, Sakura Finetek U.S.A., Inc., Torrance, CA), seven-micrometer sections cut, and sections transferred to glass slides (CryoJane, Instrumedics, Inc., St. Louis, MO). Cryosections were air dried for 30 min at room temperature, fixed for 5 min in ice-cold acetone, again air dried for 30 min at room temperature, and rehydrated in PBS for 5 min. Sections were blocked 1 hr at room temperature with 10% normal donkey serum and 30 mg/mL BSA in PBS. Primary antibodies were applied for 1 hr at room temperature using polyclonal rabbit anti-GPNMB antibody (diluted 1:200; R&D Systems Inc., Minneapolis, MN) or anti-F4/80 (diluted 1:200; Serotec Inc., Raleigh, NC). Primary antibody was removed by three washes (5 min each) in PBS and the sections were treated for 1 hr at room temperature with AlexaFluor conjugated secondary antibodies (1:200 dilution, Invitrogen-Molecular Probes, Carlsbad, CA) diluted in 1% normal donkey serum and 10 mg/mL BSA in PBS. After three washes in PBS, the sec-
Bone marrow derived dendritic cell cultures

BM derived cells obtained from D2 or D2-Gpnmb<sup>+</sup> mouse femur were cultured in the presence of GM-CSF (20 ng/ml) in 24 well plates (Costar Corp., Cambridge, MA). The nonadherent cells were removed from the culture and replated with fresh medium containing GM-CSF every alternate day. On day 10, cells were harvested and a subset replated with fresh medium containing GM-CSF every alternate day. On day 10, cells were harvested and a subset analysed by flow cytometry. In the current experiments, >95% of cells labeled positive for the mouse APC marker CD11c.

Analysis of aqueous humor IL18

Aqueous humor was collected as described previously [21]. IL18 protein level in the aqueous humor was measured using a commercially available ELISA assay following the manufacturer's recommended protocol (Bender MedSystems, San Bruno, CA). Since overall protein levels in D2 aqueous humor are increased in older mice (at 6 and 9 mo) [20], samples were compared by analyzing equal volumes of aqueous humor rather than normalizing to total protein.

Assay for immune deviation

Peritoneal exudate cells (PECs) were obtained 3 days following 2 ml intra-peritoneal injections of a 3% thioglycolate solution (Sigma-Aldrich, St. Louis, MO). Plastic adherent macrophages from these cells were used as APCs (>95% F4/80+). Cells were cultured overnight (1 x 10<sup>5</sup> well) in a 96-well plate in the presence of TGFβ2 (R&D Systems, CA, USA, 5 ng/ml final concentration) in serum free culture media, and pulsed with ovalbumin antigen (OVA) (7 mg/ml). Adherent cells were harvested, after washing off culture medium with cold Hank's balanced salt solution, and intravenously injected into B6D2F1 recipients (5 x 10<sup>3</sup> cells/mouse). Seven days following, recipients were immunized subcutaneously into the nape of the neck with OVA/CFA (50 µg). Animals that did not receive any APCs or immunization served as a negative control. A week later, animals in all groups received intradermal inoculation of OVA (200 µg/20 µl) into their right ear pinna. The left ear served as an untreated control. Thickness of both ears was measured immediately before and at 24-h interval after the OVA injection using a micrometer (Mitutoyo 227-101, MTI Corp., Paramus, NJ, USA). The measurements were performed in triplicates. Delayed type hypersensitivity (DTH) was measured as ear swelling [(24-h measurement – 0-h measurement in the experimental ear) – (24-h measurement – 0-h measurement in the untreated control ear)]. Immune deviation was detected as the suppression in DTH in the recipients of TGFβ2 treated APCs as compared to those infused with untreated APCs. A two-tailed Student's t-test was used with significance assumed at P < 0.05. DTH results were confirmed by repeating the experiments a second time.

Serum free medium

Serum-free medium (SFM) was used for in vitro assays. The medium contained: RPMI-1640, 10 mM HEPES, 0.1 mM NEAA, 1 mM Sodium pyruvate, 100 U/ml Penicillin, 100 mg/ml Streptomycin (Bio Whittaker, Walksville, MD), 0.1% BSA (Sigma-Aldrich, St. Louis, MO) and ITS+ culture supplement [1 µg/ml iron-free transferrin, 10 ng/ml linoleic acid, 0.3 ng/ml Na<sub>2</sub>Se and 0.2 µg/ml Fe(NO<sub>3</sub>)<sub>3</sub>] (Collaborative Biomedical Products, Bedford, MA).

Genetic nomenclature and stocks

The official full names of genes (with abbreviations in parentheses), alleles, and stocks utilized in this study are as follows. 1. Glycoprotein (transmembrane) nmb (Gpnmb). The ipd allele of Gpnmb results from the R150X premature stop codon mutation, Gpnmb<sup>R150X</sup> [22]. 2. Tyrosinase-related protein 1 (Tyrp1). The b allele of Tyrp1 encodes two amino acid substitutions compared to the C57BL/6J-derived allele [56]. This D2 allele is also referred to as isa in the mouse genome database and elsewhere [22,23]. The DBA/2J stock utilized here was stock number 000671 from The Jackson Laboratory. 3. Recombination activating gene 1 (Rag1). The tm1Mom mutation is a targeted knock-out [66]. The Rag1<sup>tm1Mom</sup> stock utilized here was B6.129S7-Rag1<sup>tm1Mom</sup>/J (stock 002216 from The Jackson Laboratory). 4. Protein kinase, DNA activated, catalytic polypeptide (Prkd). The scid mutation arose spontaneously in the CB17 strain and has since been backcrossed onto C57BL/6J genetic background [67]. The Prkd<sup>scid</sup> stock utilized here was B6.CB17-Prkd<sup>scid</sup>/Sjcl (stock 001913 from The Jackson Laboratory).

Authors' contributions

MGA, KSN, SM and SWMJ conceived experiments, analysed the data and prepared the manuscript. SWMJ oversaw all aspects of the study. MGA bred and performed clinical analysis of mice. KSN performed Western, ELISA
and bone marrow derived dendritic cell culture experiments. LAA and CT performed tissue sectioning and immunohistochemistry. AM performed quantitative PCR. SM performed the ACAID and quantitative PCR experiments using PECs.

Additional file 1

Loss of adaptive immune reactions has no influence on the age of onset, rate of progression, or final severity of the Tyrp1 or Gpnmb mediated iris disease. The Rag1I22041/M and Prkdcw13.6 mutations both act recessively and result in essentially identical phenotypes characterized by loss of adaptive immune responses due loss of mature B and T lymphocytes. Stocks carrying each of these mutations were separately bred to B6.Tyrp1C57BL/6GpnmbR150X mice. During the course of establishing triple mutant stocks, single and double mutant progeny classes were also isolated. Subsets of these were aged, with typical eyes of indicated genotypes shown. The combined analysis strikingly demonstrates that the immune deficiency associated with Rag1 or Prkdc mutation has no effect on Tyrp1 or Gpnmb mediated iris disease. Each column corresponds to a particular genotype with respect to Tyrp1 and Gpnmb; the rows are organized according to Rag1 or Prkdc genotype and age (homozygous mutants labeled as “Immune Deficient”, heterozygous and wild-type mice labeled “Normal”). (A to D) Eyes of young mice of all genotypes are characterized by healthy irides. By 8–9 mo, Tyrp1 and Gpnmb mutant mice exhibited typical signs of early disease that did not differ whether the mice were (E to H) immune competent, or (I to L) immune deficient. At 14+ mo, Tyrp1 and Gpnmb mutant mice exhibited signs of late disease that did not differ whether the mice were (M to P) immune competent or (Q to T) immune deficient. Panels K and Q are based on observations of 2 eyes, panel I from 6 eyes, and all other panels from groups of 8 to 26 eyes. Click here for file [http://www.biomedcentral.com/content/full/1471-2156-9-30-S1.jpeg]

Additional file 2

GpnmbR150X mutation does not directly elevate IOP. Mean IOP ± SEM is presented. For ease of comparison, some of the data are duplicated from Figure 3. It is clear that the GpnmbR150X mutation contributes to the iris disease of DBA/2J mice and is a critical component of the glaucoma. Manipulations that prevent severe iris disease in DBA/2J mice prevents the iris disease of DBA/2J mice and is a critical component of the glaucoma. Each column corresponds to a particular genotype with respect to Tyrp1 and Gpnmb; the rows are organized according to Rag1 or Prkdc genotype and age (homozygous mutants labeled as “Immune Deficient”, heterozygous and wild-type mice labeled “Normal”). (A to D) Eyes of young mice of all genotypes are characterized by healthy irides. By 8–9 mo, Tyrp1 and Gpnmb mutant mice exhibited typical signs of early disease that did not differ whether the mice were (E to H) immune competent, or (I to L) immune deficient. At 14+ mo, Tyrp1 and Gpnmb mutant mice exhibited signs of late disease that did not differ whether the mice were (M to P) immune competent or (Q to T) immune deficient. Panels K and Q are based on observations of 2 eyes, panel I from 6 eyes, and all other panels from groups of 8 to 26 eyes. Click here for file [http://www.biomedcentral.com/content/full/1471-2156-9-30-S2.jpeg]

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