IgG transmitted from allergic mothers decreases allergic sensitization in breastfed offspring

Adam P Matson1,2, Roger S Thrall1, Ektor Rafti1,3, Elizabeth G Lingenheld1 and Lynn Puddington*1

Abstract

Background: The mechanism(s) responsible for the reduced risk of allergic disease in breastfed infants are not fully understood. Using an established murine model of asthma, we demonstrated previously that resistance to allergic airway disease transmitted from allergic mothers to breastfed offspring requires maternal B cell-derived factors.

Objective: The aim of this study was to investigate the role of offspring neonatal Fc receptor for IgG uptake by intestinal epithelial cells (FcRn) in this breast milk transferred protection from allergy.

Methods: Allergic airway disease was induced during pregnancy in C57BL/6 female mice. These allergic mothers foster nursed naive FcRn+/- or FcRn-/- progeny born to FcRn+/- females that were mated to C57BL/6J-FcRn-/- male mice. In offspring deficient in FcRn, we expected reduced levels of systemic allergen-specific IgG1, a consequence of decreased absorption of maternal IgG from the lumen of the neonatal gastrointestinal tract. Using this model, we were able to investigate how breast milk IgG affected offspring responses to allergic sensitization.

Results: Levels of maternal antibodies absorbed from the breast milk of allergic foster mothers were determined in weanling FcRn-sufficient or -deficient mice. Maternal transmission of allergen-specific IgG1 to breastfed FcRn-/- offspring was at levels 103-104 lower than observed in FcRn+/- or FcRn+/+ mice. Five weeks after weaning, when offspring were 8 wk old, mice were sensitized and challenged to evaluate their susceptibility to develop allergic airway disease. Protection, indicated by reduced parameters of disease (allergen-specific IgE in serum, eosinophilic inflammation in the airways and lung) were evident in FcRn-sufficient mice nursed as neonates by allergic mothers. In contrast, FcRn-deficient mice breastfed by the same mothers acquired limited, if any, protection from development of allergen-specific IgE and associated pathology.

Conclusions: FcRn expression was a major factor in determining how breastfed offspring of allergic mothers acquired levels of systemic allergen-specific IgG1 sufficient to inhibit allergic sensitization in this model.

Background

The beneficial effects of breastfeeding on infant health have been recognized for thousands of years across diverse civilizations [1]. As breast milk is the main source of passive immunity during the early months after birth, breastfeeding is considered to be the most effective means of preventing death in young children from infectious causes [2]. In addition, breastfeeding provides nutritional, developmental, psychological, social, economic, and environmental benefits [3]. While there is overwhelming evidence supporting the role of breastfeeding in protecting children from most immune-mediated diseases [4], the components in breast milk responsible for mediating this protection are not well defined.

Maternal transfer of IgG endows offspring with short-term protective immunity [5-7]. The human fetus acquires a substantial amount of maternal IgG in utero, transported across the placenta by the neonatal Fc receptor (FcRn) [8]. In both humans and rodents, maternal IgG is acquired from breast milk [9,10], absorbed from the gut lumen via FcRn-dependent transcytosis in intestinal epithelial cells [11-14]. It is known that mice deficient in either chain of FcRn (α-chain or β2 microglobulin) have impaired capacity to absorb maternal IgG from breast milk and accelerated decay of all IgGs, but not other Ig isotypes [13,15-19]. The structure of FcRn is well charac-
terized [12,20] and several studies demonstrate a dynamic role of this receptor beyond the neonatal period [21,22].

It remains uncertain how maternal IgG acquired from breast milk impacts the susceptibility or severity of allergic diseases in children. It is known from animal models that offspring that receive serum fractions containing high titers of maternal antigen-specific IgG have suppressed IgE responses and enhanced IgG responses following immunization [23]. Similarly, the presence of maternal allergen-specific IgG1 at the time of immunization can inhibit IgE responses directed against the same allergen [24,25]. In contrast, passive transfer of allergen-specific IgG, followed by local allergen challenge within the respiratory tract can induce airway eosinophilia accompanied by hyperresponsiveness to irritants (analogous to induced bronchoconstriction in asthmatics) [26]. The effect of passive immunization on exacerbation of allergic airway disease (AAD) appears mediated by enhanced allergen uptake in airway antigen presenting cells capable of activating proinflammatory CD4+ T cells [27].

We demonstrated that the breast milk from allergic mothers can protect offspring from ovalbumin (OVA)-induced AAD; with the protective effect dependent on intact maternal B cell immunity [28]. Offspring nursed by wildtype allergic foster mothers have less severe OVA-induced AAD than offspring nursed by B cell deficient allergic foster mothers. The aim of the current study was to investigate the role of offspring FcRn in acquiring this maternal B cell-derived protective factor. We demonstrated that levels of OVA-specific IgG1 absorbed from the gut into the circulation of breastfed offspring was determined by offspring FcRn expression. Furthermore, the allergen-specific IgG1 absorbed from breast milk played a major role in preventing allergic sensitization in this model.

Methods

Animals

C57BL/6J-wildtype or FcRn-deficient (FcRn−/−) mice were obtained from Jackson Laboratories (Bar Harbor, ME) or bred in our colony at the University of CT Health Center. All mice were fed sterile food and water, and housed in microisolators under specific pathogen-free conditions. Their care was in accordance with institutional and Office of Laboratory Animal Welfare guidelines.

The generation and characteristics of FcRn−/− mice have been described [13]. For genotyping, tail pieces were obtained from mice prior to weaning and again at sacrifice. Genomic DNA was isolated using a Wizard Genomic DNA purification kit (Promega Corporation, Madison, WI) according to the manufacturer’s instructions. PCR was performed as described [13] using FcRn o393 Forward 5′-GGATGCCACTGCCCCTG-3′ and FcRn o394 Reverse 5′-CGAATTCCCCAGTGTTATT-3′primers to amplify a 248 bp fragment from the wildtype allele. Targeting vector specific o395 Forward 5′-GGAATTCCTCA GTGAAGGCC-3′ and FcRn o394 Reverse were used to amplify a 378 bp fragment from the mutant allele. Gene segments were amplified using 1 μL of purified DNA in the presence of Taq DNA Polymerase (Denville Scientific Inc., Metuchen, NJ), 2.5 mM MgCl2, 10 mM dNTPs, and 0.4 μM o393, o394, o395 primers. After 35 amplification cycles, DNA fragments were separated by electrophoresis in a 1.5% agarose gel containing ethidium bromide and visualized under ultraviolet light. FcRn+/+, FcRn+/−, and FcRn−/− mice can be distinguished using this strategy [13].

Generation of allergic airway disease (AAD)

Mice were immunized twice, separated by 7 days, by intraperitoneal injection with 0.32 μg OVA (grade V, Sigma Chemical Co., St. Louis, MO) adsorbed to 0.08 mg Al(OH)3 per gram body weight. Ten to 19 days following the second immunization, animals were exposed daily to aerosolized antigen generated from 1% OVA in normal saline with a Bioaerosol Nebulizing Generator (BANG, CH Technologies, Inc., Westwood, NJ). Exposures were 1 hour for 4 or 7 consecutive days delivered via a nose-only inhalation exposure chamber with space for exposing 48 mice simultaneously (In-Tox Products, Moriarty, NM).

Allergic mothers were generated using an adaptation of this protocol essentially as described [29]. Following 7 days of primary aerosol exposure, female mice were allowed to recover for a period of 50 days and then bred with naïve C57BL/6J male mice. Pregnant mice were subjected to a secondary challenge with aerosolized OVA daily, during embryonic days (E) 11-17 of pregnancy (duration of pregnancy in C57BL/6 mice being 19-20 days).

Sample collection for assessment of OVA-induced AAD

Severity of OVA-induced AAD was evaluated in adult mice, some of whom had been foster nursed by allergic versus naïve control mothers. Nomenclature for offspring was denoted by FcRn genotype followed by nursing mother’s immune status (see Table 1). Mice were sacrificed 24 hours after the last aerosol exposure to determine serum OVA-specific Ig concentrations, distribution of airway leukocytes, and to evaluate lung histopathology. Bronchoalveolar lavage (BAL) was performed under terminal ketamine/xylazine anesthesia. Lungs from each animal were lavaged in situ with five-1 ml aliquots of sterile saline. Numbers of total leukocytes were obtained using a Z2™ Coulter Counter (6-20 μm; Beckman Coulter, Fullerton, CA). Differential leukocyte counts were enu-
merated in BAL fluid using fluorescence flow cytometry. The live leukocyte population was identified by expression of the leukocyte common antigen CD45 [30]. CD45+ cells were analyzed by forward scatter (FSC) vs. CD11b to differentiate leukocyte subsets. Using this method, eosinophils (FSC low/CD11b high) were differentiated from macrophages (FSC high/CD11b intermediate) and lymphocytes (FSC low/CD11b negative). If neutrophils were present, they would be FSC intermediate-high/CD11b very high (our unpublished results).

For lung histology, the right lower lobe from each animal was removed, fixed with 10% formalin, processed in a standard manner, and tissue sections stained with H&E [30]. The degree of allergic lung inflammation was determined in specimens without the examiner having knowledge of the experimental condition.

**Fluorescence flow cytometry**

Monoclonal antibodies used to identify airway leukocytes were anti-CD45-FITC (30-F11), -CD11b-PerCP-Cy5.5 (M1/70), -CD19-PE (1D3), -CD8α-PerCP (53-6.7), -CD4-PE (RM4-5), -CD90.2-APC (53-2.1) purchased from BD PharMingen (San Diego, CA), and -IL33 receptor-biotin (T1/ST2, [31]) purchased from MD Biosciences (St. Paul, Minnesota). Cy5-conjugated streptavidin purchased from Jackson Immuno Research (West Grove, PA) was used to identify cells labeled with biotinylated antibodies. Cells (10⁴-10⁶) were incubated with 100 μl of appropriately diluted antibodies in PBS containing 0.2% BSA and 0.1% NaN₃ for 30 min at 4°C, and then washed with the same buffer. H-2Kb tetramer containing the OVA-derived peptide SIINFEKL was generously provided by Dr. Leo Lefrançois (University of CT Health Center, Farmington, CT) and labeling of OVA-specific CD8+ cells was as described [32]. Relative fluorescence intensities were determined on a 4-decade log scale by flow cytometric analysis using a FACSCalibur™ (Becton Dickinson, San Jose, CA).

**Determination of serum IL-5 and OVA-specific Ig levels**

In some experiments, serum was collected 24 hours after the first aerosol challenge for measurement of IL-5 levels [28]. Serum IL-5 concentrations were determined by ELISA (Pierce Biotechnology Inc., Rockford, IL). The assay was performed according to the manufacturer’s recommendation. The minimum concentration of IL-5 detectable with this assay is 1.0 pg/ml.

Serum OVA-specific Ig levels were measured by ELISA using isotype-specific capture antibodies. BD Falcon Microtest™ plates (BD Falcon, Franklin Lakes, NJ) were coated with rat anti-mouse IgG1 (A85-3), IgE (R35-72) (BD PharMingen) or goat anti-mouse IgA (Southern Biotechnology Associates), at 2 μg/ml in 0.1 M Carbonate (pH 9.5) for 16 hours at 4°C. After blocking non-specific binding, isotype-specific antibodies were captured in duplicate, as 3–4, two-fold serial dilutions of serum (within established linear ranges of the standard for each individual isotype). Detection of antigen-specific antibodies was with OVA-digoxigenin conjugates followed by anti-digoxigenin-peroxidase (Roche Diagnostics, Indianapolis, IN) [29,33]. Development was with the TMB microwell peroxidase substrate system (Kirkegaard & Perry Laboratories, Gaithersburg, MD) and A₄₅₀ measured with a Biorad Model 480 microplate reader (Hercules, CA). Limits of detection for OVA-specific IgG1, IgA, and IgE antibodies in the ELISA were 0.3 ng/ml, 10 ng/ml, or 5 ng/ml. Limits of detection in serum samples were determined by the dilution required to achieve positive readings relative to the lowest reproducible standard concentration, thus were 30, 1000, or 50 ng/ml, respectively.

**Statistical analysis**

Results are expressed as mean ± standard error of the mean (SEM). Differences in antibody levels, airway inflammatory cells, and cytokine levels between groups were determined using nonparametric Mann-Whitney or Kruskal-Wallis tests. All statistical comparisons were per-
formed with Prism 4 (GraphPad Software, San Diego, CA). Statistical significance was defined as a p value ≤ 0.05. Half life was calculated using the following formula: 
\[ t_{1/2} = \frac{\log 0.5}{\log A_e/A_0} \times t \]
where \( t_{1/2} \) is the half-life of antibody decay, \( A_e \) is the amount of antibody remaining, \( A_0 \) is original amount of antibody at day 0, and \( t \) is elapsed time [34].

**Results**

FcRn-deficient mice were susceptible to OVA-induced AAD

Prior to performing adoptive nursing studies to elucidate the role of ingested maternal allergen-specific IgG1 in protecting offspring from AAD, it was necessary to determine whether wildtype and FcRn+/- mice developed comparable parameters of allergic disease. Five to 6 week old C57BL/6J wildtype (B6) or FcRn+/- female mice were immunized with OVA adsorbed to Al(OH)3 and challenged with aerosolized OVA as described in the Methods. Serum collected 24 hours after the first aerosol challenge demonstrated equivalent levels of IL-5 in B6AAD and FcRn+/-AAD mice (Figure 1A). Following sensitization, prior to aerosol challenge, serum IL-5 in immune competent mice is minimal [28]. Similarly, during acute disease following 7 days of OVA aerosol challenge, comparable levels of OVA-specific IgG1 and IgE were present in the serum from B6AAD and FcRn+/-AAD mice (Figure 1B and 1C). Airway leukocyte populations recovered from the BAL were virtually identical with equivalent numbers of eosinophils, lymphocytes (Figure 1D and 1E), and macrophages (10-100 × 10^3 cells per mouse, data not shown) represented. In addition, similar numbers of T lymphocytes potentially participating in disease pathogenesis were recovered from the airways of B6AAD or FcRn+/-AAD mice. These T cell subsets included Th2 cells (IL-33R+CD4+) [31] and OVA-specific CD8+ cells (OVA Tetramer+CD8+) (Figure 1E). Routine histology of lung sections obtained from B6AAD or FcRn+/-AAD mice demonstrated patterns of perivascular and peribronchial inflammation, predominantly composed of lymphocytes and eosinophils, typical of the pathology we consistently observe in this model (data not shown) [28,35]. These data demonstrated that FcRn played little or no role in development of allergen-specific T and B cell responses and eosinophilic inflammation of the lungs and airways when mice were subjected to this acute model of OVA-induced AAD. Similar airway eosinophilia and histological features between FcRn-sufficient and -deficient mice using a model of mild AAD were recently reported by Nakata et al. [36].

Adoptive nursing strategy

To determine the contribution of transferred maternal allergen-specific IgG1 in the ability of breast milk from allergic mothers to protect offspring from AAD, we performed the experiment outlined in Figure 2. Naive C57BL/6J-FcRn+/- females (B6naive) were mated to C57BL/6J-FcRn+/- males, generating FcRn+/- or FcRn/- progeny. Within 24 hours of delivery, pups with or without FcRn were adoptively nursed by B6AAD foster mothers. Using this strategy where all fostered pups were born to naive mothers, acquisition of maternal allergen-specific IgG1 was restricted to breast milk. In this experiment, FcRn/- offspring were expected to have reduced systemic levels of OVA-specific IgG1 as a consequence of decreased absorption of maternal IgG from the lumen of the neonatal gastrointestinal tract [13]. Five weeks following weaning, all offspring were subjected to allergic sensitization and aerosol challenge to induce AAD as described in the Methods. Requisite controls to evaluate how acquiring allergen-specific IgG1 in breast milk affected severity of AAD were FcRn+/- pups that were born to and remained with their naïve FcRn+/- mothers (positive controls for disease) and wildtype FcRn+/- pups that were born and remained with their B6AAD mothers (positive controls for protection) [28].

Levels of OVA-specific IgG1, absorbed from breast milk of allergic mothers

FcRn+/- or FcRn/- offspring were nursed by B6AAD foster mothers using the adoptive nursing strategy (Figure 2), FcRn+/- offspring were nursed by their own B6AAD birth mothers. Sera were obtained from FcRn+/-, FcRn+/-, or FcRn/- offspring immediately prior and 4 weeks after weaning (at 24 and 52 days of life) for measurement of passively acquired maternal antibodies. As anticipated, at 24 days of life FcRn+/- and FcRn+/- offspring had similar OVA-specific IgG1 serum concentrations (14,280 ± 1861 μg/ml and 6,954 ± 1259 μg/ml respectively; Figure 3). In contrast, FcRn+/- offspring displayed significantly reduced OVA-specific IgG1 serum concentrations (< 6 μg/ml). Thus, at weaning OVA-specific IgG1 antibodies were evident in the serum of FcRn+/- offspring nursed by B6AAD mothers, however the magnitude was 10^3-10^4 lower than that observed in FcRn+/- and FcRn+/- offspring. No OVA-specific antibodies were detected in the serum of pups nursed by B6naive mothers (data not shown).

At 52 days of life (4 weeks after weaning and 1 week prior to the 1st OVA immunization), OVA-specific IgG1 concentrations were approximately 10 fold lower in FcRn+/- and FcRn+/- offspring than detected at 24 days of life (weaning). At this time, OVA-specific IgG1 antibodies were no longer detected in the serum of FcRn+/- offspring (limit of detection was 30 ng/ml based on serum dilution of 1:100). There was no difference in the t_{1/2} of ingested maternal IgG1 in serum of FcRn+/- or FcRn+/- off-
Figure 1 Similar parameters of OVA-induced AAD in wildtype or FcRn deficient mice. Five to 6 week old female C57BL/6J wildtype (B6AAD) or FcRn deficient (FcRn-/- AAD) mice were given two immunizations with OVA-Al(OH)3 followed by challenge for 7 days with 1% aerosolized OVA (daily exposure time 60 min). Parameters of disease severity measured were (A) serum IL-5 concentrations determined 24 hours after the first aerosol exposure, (B and C) serum OVA-specific Ig titers determined 24 hours after the last aerosol exposure, and (D) distribution of airway leukocytes determined by fluorescence flow cytometry as described in the Methods. Numbers of IL-33R+ or OVA-tetramer+ cells were of CD4+ and CD8+ T lymphocytes, respectively. Results expressed as means ± SEM and represent 5-6 mice per group. There were no statistical differences in disease parameters between groups. Similar parameters of disease were obtained in an independent experiment.

Figure 2 Strategy to determine the role of “offspring” FcRn in the maternal transmission of allergic protection.
ies at 52 days of life in FcRn-/- offspring, we were unable to
Given the inability to detect OVA-specific IgG1 antibod-
ies have established that IgG decay is accelerated in
IgG1 levels in individual mice at 24 days (weaning) and 52
spring (~8.5 days) when calculated from OVA-specific
the serum of pups nursed by B6naive mothers (data not
Results are presented as 12-19 individual mice per group and
we were able to calculate the t1/2 of maternal IgG1 in these mice. Previous
Given the inability to detect OVA-specific IgG1 antibod-
52 days of life was calculated to be 1.9 × 10^{-3} ng/ml, which is well below the limit of detection (30 ng/ml)
in the ELISA assay. At 52 days of life, OVA-specific IgA or
Maternal allergen-specific IgG, prevented allergic
sensitization
Adult (59 day old) mice (identified as described in Table
immunization would be < 0.2 pg/ml.
Similar concentrations of OVA-specific IgE were
observed in the serum of FcRn -/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
observed in the serum of FcRn-/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
Maternal allergen-specific IgG, prevented allergic
sensitization
Adult (59 day old) mice (identified as described in Table
immunization would be < 0.2 pg/ml.
Similar concentrations of OVA-specific IgE were
observed in the serum of FcRn -/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
observed in the serum of FcRn-/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
Maternal allergen-specific IgG, prevented allergic
sensitization
Adult (59 day old) mice (identified as described in Table
immunization would be < 0.2 pg/ml.
Similar concentrations of OVA-specific IgE were
observed in the serum of FcRn -/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
observed in the serum of FcRn-/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
Maternal allergen-specific IgG, prevented allergic
sensitization
Adult (59 day old) mice (identified as described in Table
immunization would be < 0.2 pg/ml.
Similar concentrations of OVA-specific IgE were
observed in the serum of FcRn -/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
observed in the serum of FcRn-/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
Maternal allergen-specific IgG, prevented allergic
sensitization
Adult (59 day old) mice (identified as described in Table
immunization would be < 0.2 pg/ml.
Similar concentrations of OVA-specific IgE were
observed in the serum of FcRn -/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
observed in the serum of FcRn-/-/B6AAD offspring (924
ng/ml ± 214 ng/ml) as compared to FcRn+/-/B6naive con-
Figure 3 Absorption of OVA-specific IgG, by breastfed offspring
was determined by offspring FcRn expression. Naive C57BL/6J-
FcRn+//- females (B6naive) were mated to C57BL/6J-FcRn+/+ males. Progeny of this mating were FcRn+/+ or FcRn+/-.
C57BL/6J OVA-induced AAD (B6AAD) foster mothers were generated (as described in the Methods) and within 24 hours of delivery, pups with or without FcRn were adoptively nursed by B6AAD foster mothers. Serum was collected from FcRn+/+, FcRn+/-, or FcRn-/- offspring at weaning (24 days of life) and 52
days of life (1 week prior to OVA-immunization) and concentrations of OVA-specific Igs were measured by ELISA. OVA-specific Igs were ab-
sent from the serum of pups nursed by B6naive mothers (data not
shown). Results are presented as 12-19 individual mice per group and
the red line is the mean. There were no significant differences in serum concentrations of OVA-specific IgG1 antibodies between FcRn+/- and
FcRn+/+ offspring at 24 days or 52 days of life. At 24 days of life, serum
OVA-specific IgG1 concentrations were significantly lower in FcRn -/- offspring when compared to FcRn+/- or FcRn+/+ offspring (p ≤ 0.01). At 52
days of life, OVA-specific IgG1 antibodies were no longer detected in
the serum of FcRn-/- offspring (limit of detection 30 ng/ml).
Figure 4B), mononuclear cells (266 ± 41 × 10^3, data not shown), and lymphocyte subsets (Figure 4C). Furthermore, histopathologic examination of lung tissue obtained from FcRn−/−/B6AAD offspring demonstrated extensive perivascular and peribronchiolar cuffing, and eosinophilic inflammation, similar to that previously reported for wildtype C57BL/6 mice in this model (Figure 4D) [35]. In contrast, there was notably less allergic inflammation in lung tissue obtained from FcRn+/+/B6AAD and FcRn+−/−/B6AAD offspring (Figure 4D). Thus, consistent with our previous study [28], FcRn-sufficient offspring nursed by wildtype OVA-immune mothers were protected from developing severe OVA-induced AAD. In contrast, limited, if any, protection from development of OVA-induced AAD was transmitted from allergic mothers to FcRn-deficient offspring.

**Discussion**

There is overwhelming evidence supporting the role of breastfeeding in protecting children from most immune-mediated diseases [4]. Despite this, it is not clear whether this applies to prevention of allergic disease in situations when mothers are allergic. Possible explanations for the inconsistent effects of breastfeeding on allergy and asthma prevention may be the immunologic complexities of breast milk itself and potential changes in composition in the context of maternal allergy or allergen exposure. Breast milk contains a multitude of biologically active components and some elements are thought to protect the infant from developing allergies, whereas others might promote allergic sensitization [37].

We recently demonstrated that transmission of resistance to AAD from allergic mothers to nursing offspring is dependent on B cell-derived factors in breast milk [28]. By comparing offspring of mothers with OVA- versus BSA-induced lung disease, we also established that the maternally transferred protection from AAD is antigen-specific [29]. Based on these findings, we hypothesized that antigen-specific IgG in breast milk were major contributors to this protective effect. We previously demonstrated that allergen-specific IgG, IgA and IgE are absorbed from the neonatal gastrointestinal tract into the systemic circulation of naïve mice nursed by allergic mothers [28,29]. No allergen-specific IgG2a is elicited following immunization with OVA adsorbed to Al(OH)₃ or after the aerosol challenge in our model of AAD, thus is not absorbed by offspring nursed by allergic mothers [29]. In the present study, although maternal allergen-specific IgG₁, IgA and IgE were present at weaning in naïve FcRn-sufficient mice foster nursed by allergic mothers, it appeared that allergen-specific IgG₁ was the
only isotype whose levels were sustained until allergic sensitization. This was not the case in FcRn-deficient mice where the low levels of maternal allergen-specific IgG1 present at weaning were undetectable at the time of immunization. In mice, as in humans, the transfer of breast milk IgG across the intestinal epithelium is mediated by FcRn [9]. Based on these and other data presented in this report, we were able to show that expression of FcRn was important for offspring to acquire sufficient levels of allergen-specific IgG1 from the breast milk of allergic mothers to prevent allergen-specific IgE responses. A distinct experimental strategy to address this research question was recently reported by Nakata et al. [36]. Importantly, the data derived from their studies led them to the same overall conclusion, that maternal IgG affects development of allergy in offspring. Thus, the two studies synergize to advance the understanding of the biology of FcRn as it applies to uptake of maternal IgG from the lumen of the gastrointestinal tract, and of how absorbed maternal allergen-specific IgG and offspring FcRn contribute to enhancing protection from allergic sensitization and disease.

To determine the role of FcRn in the postnatal acquisition of allergic protection, it was necessary to establish that FcRn-deficient mice were competent to develop OVA-induced AAD. After OVA-immunization and aerosol challenge, FcRn-/- AAD mice demonstrated equivalent parameters of acute disease as wildtype B6AAD mice (this report and [36]). Of particular interest were similar titers of OVA-specific IgG1 antibodies in FcRn-/- AAD and B6AAD mice in serum collected 24 hours after the last aerosol exposure. Thus, despite the accelerated decay of IgG in FcRn-deficient mice [13,15,16], the initial antibody titers following aerosol challenge were unaffected. These results suggest that lymphocyte responses to allergic sensitization and challenge, including the generation of OVA-specific memory CD4+ T cells and B cells, were intact in FcRn-deficient mice. Furthermore, differentiation of memory B cells to antibody-producing plasma cells appeared unaffected by the absence of FcRn.

FcRn-/- offspring had impaired capacity to absorb OVA-specific IgG1 from the breast milk of allergic mothers. At 24 days of life, 10^3 - 10^4 lower levels of antigen-specific IgG1 were detected in the serum of FcRn-/- offspring as compared to FcRn+/- or FcRn+/+ offspring nurtured under the same conditions. This is consistent with what is known regarding the significant role of FcRn in mediating absorption of breast milk IgG [13,17]. However, in the previous study, while the TNP-specific IgG1, injected into pregnant mice was present in the serum of breastfed FcRn+/- neonates (10-20 μg/ml), it was not detected (<80 ng/ml) in their littermate FcRn-/- mice [13]. Thus, the existence or impact of an FcRn-independent component of maternal IgG uptake has not been appreciated. It is possible the low levels of antigen-specific IgG1 detected in the serum of FcRn-/- offspring are acquired via passive diffusion across the intestinal epithelium, although it remains to be determined whether this is the case. It is known that this mucosal barrier is more permeable in neonates with gut "closure" (cessation of Ig absorption) occurring at weaning [10].

Although we demonstrated that FcRn-independent uptake of maternal IgG can occur in neonatal mice, we found that >99.9% of IgG absorbed in wildtype mice was via an FcRn-dependent mechanism (see Figure 3). In addition to mediating transcytosis of IgG across the intestinal epithelium [11-13], FcRn modulates IgG homeostasis [13,15,16]. Thus, we were able to perform the first study that quantified the rate of decay of absorbed maternal IgG1 acquired exclusively from breast milk (~8.5 days). As compared to IgG acquired via intravenous injection, it is possible that ingested IgG selected by FcRn for absorption from the gut lumen has a higher binding affinity for FcRn systemically, and thereby has increased protection from catabolism. It is not clear whether or not this is the case since a direct comparison of half-lives of the same population of IgG1 molecules following ingestion or injection of mice at the same age has not been made. Furthermore, there appears to be a lack of consensus in the field regarding the t1/2 of injected IgG [13,15,16,18,19]. The explanation for the diversity in results from different laboratories is not obvious, but could be due to different routes of injection - intraperitoneal versus intravenous, or structural features of the injected monoclonal antibodies tested that contribute to their inherent strengths of interaction with FcRn.

Other studies demonstrate that under the appropriate experimental conditions, breast milk may be protective against [28,36,38-41] or increase susceptibility to [42] the development of allergic disease in offspring. It is important to understand the mechanistic basis for differences in the effect of breast milk during this early period of immune maturation. In our studies, it is possible that maternal antigen-specific IgG1 absorbed into the systemic circulation of offspring, neutralized the antigen-clearing it from the circulation prior to its recognition by cells of the adaptive immune system. This is supported by data from offspring nursed by allergic mothers where few, if any, FcRn-sufficient offspring produced allergen-specific IgE at 7 days after the second intraperitoneal immunization (data not shown). Neutralization of pathogens is known to occur when infants receive certain live vaccines (e.g. measles virus) in the presence of preexisting maternal antibodies. This is a major factor for delaying infant immunization until the majority of maternal antibodies have disappeared [43,44]. The functions of maternal anti-
bodies in determining immune parameters in offspring can be influenced by the presence or absence of antigen and the ratio between them [45]. Interestingly, in some experiments performed in this and our other related studies evaluating how allergic mothers influence parameters of allergic disease in offspring, allergen-specific IgG1 and IgE responses were virtually absent without profound effects on airway eosinophilia. Since airway eosinophilia can occur in the complete absence of B cells [28], this implies that protection from AAD acquired from allergic mothers in our model has more robust downstream effects on B cell than T cell parameters of disease. Thus, in addition to allergen neutralization, it is likely that maternal antigen-specific IgG1/allergen immune complexes contribute to determining the outcome of offspring responses to allergic sensitization [46].

It should be noted that not all maternal Igs have beneficial effects in progeny. Recently, a murine model of peanut allergy demonstrated that maternal derived anti-peanut IgG1 is associated with anaphylactic reactions in offspring [47]. In addition, several autoimmune diseases such as systemic lupus erythematosus are known to result in transmission of maternal IgG’s that have deleterious effects in progeny [48]. The ability of maternal IgG to mediate differential effects in offspring may be related to affinities for individual FcyRs resulting from Fc glycosylation. IgG is known to contain a single N-linked glycan at Asn297 of the Fc domain, and variations of this covalently linked complex carbohydrate determines binding interactions with individual FcyRs [49,50]. Fc sialylation of IgG1 results in a reduced binding affinity for the activating receptor FcγRIIb and promotes anti-inflammatory effects through the inhibitory receptor FcγRIIB [49]. It is known that several autoimmune diseases are associated with individual glycoforms of IgG [51,52]. Perhaps the structure of glycans on antigen-specific IgGs varies during the pathogenesis of allergic disease, or is influenced by the environment at the site of the allergen challenge (such as the lung or gut mucosa). Control of post-translational modification of carbohydrate residues on IgGs could be determined during plasma cell differentiation from memory B cells, and/or modified by glycosylation or de-glycosylation enzymes unique to specific tissue environments or disease states. It is possible that physiological changes during lactation itself play a role in the characteristics of maternal IgG acquired by offspring to influence whether maternal IgG transfers increased risk or protection from allergic disease. Recent results from Victor et al. [41] could be supportive of this idea. In their study, neonates nursed by immunized mothers exhibit marked inhibition of B and T cell responses following immunization. In contrast, postnatal injected anti-allergen IgG (purified from serum of immunized mice) failed to modulate expression of FcγRIIB or regulate B or T cell cytokine production.

Our findings suggest a serum concentration limit of absorbed maternal antigen-specific IgG sufficient to protect offspring from AAD. This concept is supported by data from FcRn+/+ or FcRn−/− offspring, where serum levels of antigen-specific IgG1 of 10^3 - 10^6 ng/ml at 52 days of life appeared sufficient to protect offspring from AAD initiated one week later. Significantly reduced levels of antigen-specific IgG1 in the serum of weaning FcRn−/− mice, that decayed to negligible levels prior to immunization, resulted in the absence of protection from AAD. Interestingly, the concentration limit of maternal IgG needed to protect offspring from AAD appears to be dependent on the severity of disease elicited in murine models, with lower levels of absorbed allergen-specific IgG1 (60-90 ng/ml) being sufficient to protect from mild disease [36]. Additional experimentation aimed at defining the contributions of serum concentrations of maternal IgG, immune complexes and structural glycoforms sufficient to protect offspring from allergic sensitization will be important.

Conclusion

Our study demonstrates that breast milk factors obtained via FcRn (e.g. IgG) result in reduced severity of allergic airway disease in offspring. Based on these results one could consider increasing maternal antigen-specific IgG levels (e.g. maternal immunization) as a possible method for the prevention of allergic disease in progeny. Further clarification of the IgG levels required to protect offspring, the structural properties of antibodies involved, and their interaction with receptors at various locations (e.g. spleen, intestine, and thymus) in the neonate are important in understanding how passive immunity influences the development of allergy in offspring.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

APM supervised the animal experiments, participated in the immunoassays and study design, performed the statistical analysis, and drafted the manuscript. RST reviewed the histopathology and helped to draft the manuscript. ER performed the majority of immunoassays, PCR, and sample collections. EGL helped to direct the animal experiments and immunoassays. LP conceived of the study, participated in its design, coordination, and data analysis; and helped to draft the manuscript. All authors have read and approved the final manuscript.

Acknowledgements

We are indebted to Derry Roopenian for his assistance in obtaining stocks of FcRn-deficient mice used to establish our colony. We thank Rick Blumberg and Neil Simister for their helpful discussions as we undertook this project. We are grateful to Li Zhu for her assistance in the laboratory and Eric Secor for his assistance with the photomicrographs. We deeply appreciate Michelle Cloutier and Vic Herson’s encouragement and support. This work was supported by the National Institutes of Health: KO8AI071918 (to APM) and HL080508 (to LP), and...
in part by the Burr Curtis Research Endowment, Connecticut Children’s Medi-
cal Center (to APM).

Author Details
1Department of Immunology, University of Connecticut Health Center,
Farmington, Connecticut, USA; 2Department of Pediatrics, Connecticut
Children’s Medical Center, Hartford, Connecticut, USA and 3Department of
Research, Connecticut Children’s Medical Center, Hartford, Connecticut, USA

Received: 28 April 2010 Accepted: 13 July 2010
Published: 13 July 2010

References
1. Newburg DS: Bioactive components of human milk: evolution, efficiency, and protection. Adv Exp Med Biol 2001, 501:3–10.
2. Labbok MH, Clark D, Goldman AS: Breastfeeding: maintaining an
irreplaceable immunological resource. Nat Rev Immunol 2004, 4:565-572.
3. Gartner LM, Morton J, Lawrence RA, Naylor AJ, O’Hare D, Schanler RJ, et al.
4. IgP S, Chung M, Raman G, Chew P, Magula N, Devine D, et al.
5. Ehrlich P: "Collected Papers, Vol II. Z Hygiene 1892, 12:31-44.
6. Brambell FW, Halliday R, Brierley J, Hemmings WA: Transference of
passive immunity from mother to young. Lancet 1954, 266:964-965.
7. Zinkernagel RM: Maternal antibodies, childhood infections, and
autoimmune diseases. N Engl J Med 2001, 345:1331-1339.
8. Simister NE, Story CM, Chen HL, Hunt JS: The MHC class I-like IgG receptor
controls perinatal IgG transport, serum half-lives of IgG in beta 2-microglobulin-deficient mice. J Immunol 2003, 170:3528-3533.
9. He W, Ladinsky MS, Huey-Tubman KE, Jensen GJ, McIntosh JR, Bjorkman PJ: FcRn-mediated antibody transport across epithelial cells revealed by electron tomography. Nature 2008, 455:542-546.
10. Ghette V, Hubbard IG, Kim JK, Tien MF, Lee Y, Ward ES: Abnormally short
half-life of IgG in beta 2-microglobulin-deficient mice. Eur J Immunol 1996, 26:1527-1531.
11. Jones EA, Waldmann TA: The mechanism of intestinal uptake and
transcellular transport of IgG in the neonatal rat. J Clin Invest 1972, 51:2916-2927.
12. Rodevald R, Krachenbuhl JP: Receptor-mediated transport of IgG. J Cell Biol 1984, 99:1595-164.
13. Israel EJ, Patel V, Taylor SF, Marshak-Rothstein A, Simister NE: The role of breast-feeding in the development
independent of interleukin 4, interleukin 5, and interleukin 10, and
important for Th2 effector function. Proc Natl Acad Sci USA 1998, 95:6930-6935.
14. Masopust D, Veys V, Marzo AL, Lefrançois L: Preferential localization of
effecter memory cells in nonlymphoid tissue. Science 2001, 291:2413-2417.
15. Seymour BPM, Genthin-LJ, Coffman RL: Aerosol-induced
immunoglobulin (Ig)-E unresponsiveness to ovalbumin does not require CD8+ or CD4+ T cells, or interferon (IFN)-γ, or CD40 ligand in a murine model of allergen sensitization. J Exp Med 1998, 187:271-31.
16. Roopenian DC, Christianson GJ, Sproule TJ, Human FcRn transgenic mice for pharmacokinetic evaluation of therapeutic antibodies. Methods Mol Biol 2010, 620:293-104.
17. Schramm CM, Puddington L, Whiteley HE, Yamaouyannis CA, Schramm CM, Mohrmanul F, et al: Murine cytomegalovirus infection alters Th1/Th2
cytokine expression, decreases airway eosinophilia, and enhances mucus production in allergic airway disease. J Immunol 2001, 167:2798-2807.
18. Lohning M, Stroehmann A, Coyle AJ, Grogan JL, Lin S, Gutierrez-Ramos JC, et al: T1/T2 is preferentially expressed on murine Th2 cells, independent of interleukin 4, interleukin 5, and interleukin 10, and
important for Th2 effector function. Proc Natl Acad Sci USA 1998, 95:6930-6935.
19. Matson AP, Zhu L, Lingenheld EG, Schramm CM, Clark RB, Selander DM, et al: Maternal transmission of resistance to development of allergic
airway disease. J Immunol 2007, 179:1282-1291.
20. Wu CA, Puddington L, Whiteley HE, Yamaouyannis CA, Schramm CM, Mohrmanul F, et al: Murine cytomegalovirus infection alters Th1/Th2
cytokine expression, decreases airway eosinophilia, and enhances mucus production in allergic airway disease. J Immunol 2001, 167:2798-2807.
21. Lohning M, Stroehmann A, Coyle AJ, Grogan JL, Lin S, Gutierrez-Ramos JC, et al: T1/T2 is preferentially expressed on murine Th2 cells, independent of interleukin 4, interleukin 5, and interleukin 10, and
important for Th2 effector function. Proc Natl Acad Sci USA 1998, 95:6930-6935.
22. Kobayashi K, Qiao SW, Yoshida M, Baker K, Lencer WI, Blumberg RS: An
FcRn-dependent role for anti-flagellin immunoglobulin G in pathogenesis of colitis in mice. Gastroenterology 2009, 137:1746-1756.
23. Jarrett EE, Hall E: IgE suppression by maternal IgG. Immunology 1983, 48:49-58.
24. Victor JR Jr, Fusaro AE, Duarte AJ, Sato MN: Preconception maternal
immunization to dust mite inhibits the type I hypersensitivity response of
offspring. J Allergy Clin Immunol 2003, 111:269-271.
25. Littih H, Spennier A, Reckelkamp W, Ahrens B, Wolk G, Hackler R, et al: Critical role of preconceptional immunization for protective and
nonpathological specific immunity in murine neonates. J Immunol 2003, 171:3485-3492.
26. Oshiba A, Hamelmann E, Takeda K, Bradley KL, Loader JE, Larsen GL, et al: Passive transfer of immediate hypersensitivity and allergic
hypersensitiveness by allergen-specific immunoglobulin (Ig) E and
IgG in mice. J Clin Invest 1996, 97:1398-1406.
27. von Garnier C, Wikstrom ME, Zosky G, Turner DJ, Sly PD, Smith M, et al: Allergic airways disease develops after an increase in allergen capture and
processing in the airway mucosa. J Immunol 2007, 179:5748-5759.
28. Matson AP, Thrall RS, Rathi F, Puddington L: Breastmilk from allergic
mothers can protect offspring from allergic airway inflammation. Breastfed Med 2009, 4:167-186.
29. Matson AP, Zhu L, Lingenheld EG, Schramm CM, Clark RB, Selander DM, et al: Maternal transmission of resistance to development of allergic
airway disease. J Immunol 2007, 179:1282-1291.
30. Wu CA, Puddington L, Whiteley HE, Yamaouyannis CA, Schramm CM, Mohrmanul F, et al: Murine cytomegalovirus infection alters Th1/Th2
cytokine expression, decreases airway eosinophilia, and enhances mucus production in allergic airway disease. J Immunol 2001,
167:2798-2807.
43. Siegrist CA: Neonatal and early life vaccinology. Vaccine 2001, 19:3331-3346.
44. Albrecht P, Ennis FA, Saltzman EJ, Krugman S: Persistence of maternal antibody in infants beyond 12 months: mechanism of measles vaccine failure. J Pediatr 1977, 91:715-718.
45. Lambert PH, Liu M, Siegrist CA: Can successful vaccines teach us how to induce efficient protective immune responses? Nat Med 2005, 11:554-562.
46. Mosconi E, Rekima A, Seitz-Polski B, Kanda A, Fleury S, Tissandie E, et al: Breast milk immune complexes are potent inducers of oral tolerance in neonates and prevent asthma development. Mucosal Immunol 2010, 3: doi:10.1038/mi.2010.23
47. Lopez-Exposito I, Song Y, Jarvinen KM, Srivastava K, Li XM: Maternal peanut exposure during pregnancy and lactation reduces peanut allergy risk in offspring. J Allergy Clin Immunol 2009, 124:1039-1046.
48. Tincani A, Rebaioli CB, Frassi M, Taglietti M, Gorla R, Cavazzana L, et al: Pregnancy and autoimmunity: maternal treatment and maternal disease influence on pregnancy outcome. Autoimmun Rev 2005, 4:423-428.
49. Kaneko Y, Nimmerjahn F, Ravetch JV: Anti-inflammatory activity of immunoglobulin G resulting from Fc sialylation. Science 2006, 313:670-673.
50. Nimmerjahn F, Ravetch JV: Divergent immunoglobulin G subclass activity through selective Fc receptor binding. Science 2005, 310:1510-1512.
51. Parekh RB, Dwek RA, Sutton BJ, Fernandes DL, Leung A, Stanworth D, et al: Association of rheumatoid arthritis and primary osteoarthritis with changes in the glycosylation pattern of total serum IgG. Nature 1985, 316:452-457.
52. Holland M, Yagi H, Takahashi N, Kato K, Savage CO, Goodall DM, et al: Differential glycosylation of polyclonal IgG, IgG-Fc and IgG-Fab isolated from the sera of patients with ANCA-associated systemic vasculitis. Biochim Biophys Acta 2006, 1760:669-677.

doi: 10.1186/1476-7961-8-9
Cite this article as: Matson et al. IgG transmitted from allergic mothers decreases allergic sensitization in breastfed offspring Clinical and Molecular Allergy 2010, 8:9