On the Perplexingly Low Rate of Transport of IgG2 across the Human Placenta

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Abstract
The neonatal receptor, FcRn, mediates both serum half-life extension as well as active transport of maternal IgG to the fetus during pregnancy. Therefore, transport efficiency and half-life go hand-in-hand. However, while the half-life of the human IgG2 subclass is comparable to IgG1, the placental transport of IgG2 is not, with the neonatal IgG1 levels generally exceeding maternal levels at birth, but not for IgG2. We hypothesized that the unique short-hinged structure of IgG2, which enables its k chain, but not λ isotype to form at least three different structural isoforms, might be a contributing factor to these differences. To investigate whether there was any preference for either light chain, we measured placental transport of IgG subclasses as well as k/λ-light chain isotypes of IgG1 and IgG2 in 27 matched mother-child pairs. We also studied the half-life of IgG1 and IgG2 light chain isoforms in mice, as well as that of synthesized IgG2 structural isoforms kA and kB. In order to investigate serum clearance of IgG1 and IgG2 light chain isoforms in humans, we quantified the relative proportions of IgG1 and IgG2 light chains in hypogammaglobulinemia patients four weeks after IVlg infusion and compared to the original IVlg composition. None of our results indicate any light chain preference in either of the FcRn mediated mechanisms; half-life extension or maternal transport.

Introduction
Immunoglobulin G (IgG) forms the backbone of our circulating, adaptive immune system. The fully assembled IgG molecule consists of two identical 50 kDa heavy chains (γ1, γ2, γ3 or γ4 subclasses), and two identical 23 kDa light chains forming a heterodimer (one heavy chain and one light chain) that further assemble into dimers. The assembled molecule is Y shaped, with the light chains and the N-terminal parts of the heavy chains (CH1 and VH domains) in tight association, forming the two Fab arms (Fragment antigen binding), and the C-terminal CH2 and CH3 domains forming the Fc tail. The CH1 and CH2 domains are connected by a flexible hinge, allowing the Fab ‑ C2 considerable freedom of movement from the Fc portion. Length and flexibility of the hinge region varies extensively amongst the IgG subclasses influencing the relative orientation and movement of the Fab arms and Fc tail of the IgG antibody [1].

The hinge region of IgG1 encompasses 15 amino acids and is very flexible. IgG2 has a 12 amino acid hinge region and contains a rigid poly-proline double helix, stabilized by four inter-heavy chain disulfide bridges. IgG3 has the longest hinge region, about 4 times as long as IgG1, and thus the greatest flexibility, while the IgG4 hinge contains 12 amino acids yielding an intermediate flexibility compared to IgG1 and IgG2 [2].

Light chains come in two classes, either κ or λ, with four highly homologous λ light chain allotypes. In humans the κ/λ ratio in serum is around 2:1 in healthy individuals, but this varies between species, isotype, biological location and age [3]. No functional differences between λ and κ antibodies have been described so far. Recently, IgG2κ was described to occur in three distinct isoforms, A, A/B and B, which differ from each other solely in their disulfide bridges in the hinge, with four disulfide bonds connecting the Fc chains for A, but two in the B form, and an hybrid A/B form with three inter-Fc bonds [4]. This affects its tertiary structure and thus the position and mobility of F(ab')2, which in turn may affect other interactions [5]. In contrast, IgG2κ is found predominantly as the A and A/B molecular form, but devoid of the B form [4,6].

FcRn, the neonatal FcγR, is a heterodimer of a unique MHC class I-like alpha chain and β2m. It is thought to, amongst other functions, be responsible for both the long half-life of IgG in vivo and to mediate IgG transcytosis, for instance to the mucosa (e.g.
IgG2 hinge, which causes the F(ab)2 binding affinity to mouse FcRn [23]. Due to the shortness of the humans, the IgG2 light chain isotype has been found to influence been reported to have unusually short half-life in mice [15,19–22]. The interaction still allows for half-life extension of human IgG1, IgG3 or IgG4. All human IgG subclasses have levels. Thus the transport of IgG2 is much less efficient and is in fact comparable to that of R435-containing IgG3 [17,19]. Since IgG across the placenta however, the concentration of all the histidine at 435 (H435) [13]. IgG1, IgG2 and IgG4 share all currently described contact residues with FcRn, including H435, with similar affinity to FcRn when measured using immobilized human FcRn on a biosensor and a comparable half-life in circulation [14–16]. For transport of IgG, Montaño and Morrison reported a difference in half-life between human-mouse chimeras of IgG2α and IgG2κ, with IgG2κ being cleared faster in mice [15,19]. This light-chain difference was not observed for IgG1, IgG3 or IgG4. All human IgG subclasses have been reported to display pH-dependent binding activities to mouse FcRn, albeit slightly altered compared to binding to human FcRn [16]. The interaction still allows for half-life extension of human IgGκ by mouse FcRn, with comparable relative half-lives reported, as in humans (IgG1κ≈IgG2κ≈IgG3κ), except for IgG4, which has been reported to have unusually short half-life in mice [15,19–22]. Although no difference in IgG2 half-life has been described in humans, the IgG2 light chain isotype has been found to influence binding affinity to mouse FcRn [23]. Due to the shortness of the IgG2 hinge, which causes the F(ab)2 to be relatively close to the Fc as well as the distinct structural isoforms of IgG2, we hypothesized that the low placental transport of IgG2 may be attributable to a single structural isoform, unique to IgG2κ, interacting differently with FcRn, possibly affecting its half-life as well.

For this reason we studied the trans-placental transport of IgG2α and IgG2κ in humans and their half-life in mice and humans, both of which are FcRn-ascribed functions, and tested whether the unique structural isoforms of IgG2κ could explain the low efficiency of IgG2 transport.

**Materials and Methods**

**Recombinant IgG**

V4-μ-matched recombinant IgG1 (λ and κ) and IgG2 (λ and κ) described before [24,25], were produced in 293 Freestyle cells (Invitrogen, Carlsbad CA) according to the manufacturer’s instructions with p20, p27 and SV40 Large-T antigen as described, and purified by protein G HiTrap columns using the Acta Prime Plus system (GE Healthcare, Buckinghamshire, UK) [26].

**IgG quantification**

Light chain and IgG1 and IgG2 subclass specific antibody quantification on the paired mother-cord samples was done by sandwich ELISA using Nunc MaxiSorp plates (Sigma-Aldrich, St. Louis, MO). For IgG total ELISA, mouse anti-human IgG (M1268, Sanquin) was used for coating (1/500), and HRP-mouse anti-human Fc (JDC10 Southern Biotech) was used as a secondary antibody (1/1000). For IgG light chain ELISA’s mouse-anti-human IgG1 (MH1325, Sanquin, 1/100) was used to coat, using either HRP-labelled mouse-anti-human λ (JDC12, Southern Biotech, 1/1600) or κ (HP6062, Southern Biotech, 1/1300). Similarly, mouse anti-human IgG2 (HP6002, Southern Biotech, 1/100 for lambda assay, 1/50 for kappa assay) was used for coating, and HRP-labelled mouse anti-human λ (JDC12 (Southern Biotech, 1/1600) or κ (HP6062, 1/1300) for detection.

Conversion of 3,3′,5,5′-Tetramethylbenzidine (TMB) was used to quantify HRP activity per well and absorptions were read using a Genios Pro plate reader (Tecan, Mannefedorf, Switzerland) using standard sets of filters of 450 nm.

IgG subclass concentrations in sera were determined by nephelometry (Behringer nephelometer II, Behringer diagnostics).

**In vivo serum persistence experiments**

BALB/c mice were obtained from the NKI institute in Amsterdam. C57Bl/6 and C57Bl/6-FcRn-KO mice were obtained from the Charles River and Jackson laboratories, respectively. All in vivo experiments were performed using female 8–9 week old mice weighing between 16.8 and 26.3 grams each. Each mouse received 200 μg purified recombinant human IgG of the appropriate heavy and light chain subclass and isofrom by IV administration. Blood samples of 100 μl were drawn from the tail at day 0 and then approximately every 5 days for a period of two weeks, and human IgG concentration was determined with a mouse anti-human IgG ELISA. All animal experiments were carried out after approval from the ethical committee of Sanquin/Dutch Cancer Institute (Dierexperimentencommissie NKI), who also monitored the progress, aimed at minimizing suffering (number 10.033).

In humans IgG light chain subclass serum persistence was calculated as (%) = [IgGκ]t/([IgGκ]t0)/(IgGκ]t/([IgGκ]t0) x100% in sera from three agammaglobulinemic patients receiving IVIg at 4 weeks interval as described in [20].

**IgG2κ redox treatment**

Enrichment of IgG2 κ isoforms by redox treatment was performed as described in Dillon et al 2008 (Structural and Functional Characterization of Disulfide Isoforms of the Human
IgG2 Subclass) [5]. Shortly, for the synthesis of the B isoform, IgG2k was incubated at 3 mg/ml in 200 mM Tris buffer at pH 8 with 6 and 1 mM of cysteine and cystamine, respectively. For IgG2k A enrichment, 0.9M guanidine hydrochloride (GuHCl) was also added. The samples were protected from light and placed at 2–8°C for 48–72 h. Afterwards the antibody was run through a Zeba spin desalting column (Pierce) for buffer exchange into PBS, and stored at 220°C.

SDS gel electrophoresis

5 μg of protein was loaded on a Thermo Scientific Tris Hepes SDS precast polyacrylamide mini gel 12%. Reduced samples were diluted with NuPAGE LDS Sample Buffer (4×) with 0.1% Beta MercaptoEthanol and incubated at 95°C for 5 minutes. Non-reduced samples were diluted with NuPAGE LDS Sample Buffer (4×) with 60 mM Iodoacetamide and incubated at 70°C for 5 minutes. The running buffer was TRIS HEPES SDS buffer for both types of samples, for the non-reduced samples 4 mM Iodoacetamide was added. Run time was ca 1.5 hour at 130 mA/gel, 120 V. Afterwards, gels were stained with Coomassie blue.

Surface plasmon resonance

Surface plasmon resonance was conducted using a Biacore 3000 instrument (GE Healthcare) with CM5 sensor chips. The coupling was performed by injecting 5 μg/ml of the each protein into 10 mM sodium acetate pH 4.5 using the amine coupling kit (GE Healthcare). Titrated amounts (8,000.0-62.50 nM) of monomeric human FcRn were injected over immobilized IgG variants (500 RU) using phosphate buffer (67 mM phosphate buffer, 0.15 M

Figure 1. Placental transport of IgG subclasses is more efficient for IgG1 and IgG4 than for IgG2 and IgG3. Blood was collected from mothers just before or after birth and from neonates birth. A) Transport rates for all IgG subclasses expressed as cord/maternal ratios found at birth. The transport rates differed significantly from each other (P<0.0001), except for IgG2 and IgG3 (not significant), as tested by one-way Anova and Tukey's multiple comparison test. (B–D) IgG subclass 1–4 serum levels were quantified by nephelometry and each pair was plotted on a X axis displaying days of each pregnancy against IgG concentration. Average neonate concentration was significantly higher than in the mother for IgG1 and IgG4 as tested by a paired-T test as shown (child/mother ratio = 1.55 and 1.38, respectively) while average concentrations for IgG2 and IgG3 were not significantly different in mothers and their children (child/mother ratios not significantly different from 1). One pre-term baby was identified displaying low transport of all IgG (square symbol). (F) Child/mother transport ratio of subclasses IgG2-4 for each pair was plotted relative to the IgG1 transport ratios.

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NaCl, 0.005% Tween 20) at pH 6.0 as running buffer whereas HBS-P buffer at pH 7.4 was used for regeneration of the flow cells. Experiments were conducted at 25°C with a flow rate of 40 μl/min. All sensorgrams were zero adjusted and reference cell measurements at this range, this value was excluded from further analysis.

**Results**

**IgG2 and IgG3 are not efficiently transported across the placenta**

We quantified all four IgG subclasses in 27 paired mother cord plasma samples taken at birth (Fig. 1A–D). As expected, IgG1 and IgG4 levels in cord blood were generally found to be higher than in maternal blood. Specifically, cord exceeded maternal levels in 24 out of 27 samples with the average cord/mother ratio being 1.5 for IgG1, 1.3 for IgG4 while for IgG3 the average cord/mother ratio was 1.00. IgG4 levels in cord and maternal placental plasma were also about equal (with average cord/mother ratio being 0.99). Interestingly, the pairs show a highly varied transport, with some individuals transporting all subclasses at higher efficiency than others. However, the overall transport was relatively uniform for all the subclasses: if relatively high transport was observed for IgG1, the same held true for IgG2 (R² = 0.875, p < 0.0001), IgG3 (R² = 0.836, p < 0.0001) and IgG4 (R² = 0.303, p = 0.0052) (Fig. 1E). One case with extremely strong IgG4 transport, with maternal levels being very low (0.03 g/L) but rising to 0.11 g/L in (Fig. 1E). One case with extremely strong IgG4 transport, with maternal levels being very low (0.03 g/L) but rising to 0.11 g/L in the fetal circulation (Fig. 1E). Due to the uncertainty of measurements at this range, this value was excluded from further analysis.

In general, the relative transport of all subclasses was in direct relationship to the maternal levels: the higher the maternal IgG, the lower the transport observed for all IgG subclasses (Fig. 2). The placental transport rates were IgG1 > IgG4 > IgG2 = IgG3, at all levels for maternal IgG. The slopes of the respective trend lines were IgG1: -3.664 ± 1.023; IgG2: -5.332 ± 1.861; IgG3: -4.434 ± 1.721; IgG4: -3.963 ± 1.420.

**IgG2k and IgG2λ are transported equally well across the placenta**

Montano and Morrison described that FcRn-mediated half-life of human-mouse chimeric IgG2k was shorter than IgG2k in Balb/c mice, while the FcRn-mediated half-life of light chain isotype of the other subclasses was unaffected [15]. Later the same group described increased affinity of mouse FcRn to human IgG2λ compared to IgG2k [23]. We therefore investigated if this was also true for human FcRn by measuring the affinity of soluble monomeric human FcRn to immobilized IgG2k variants at pH 6.0 using surface plasmon resonance. We found FcRn to bind IgG2k slightly stronger than IgG2k (KD = 1.1 μM and 1.7 μM, respectively) (Fig. 3). However, no difference in binding affinity was detected for the IgG1-light chain isotypes (KD = 1.3 μM).

Although somewhat counterintuitive, these findings raised the possibility that placental transport of IgG2k and λ may also differ as both half-life and placental transport is mediated by FcRn. To test this, we first set up light chain-specific IgG1k and IgG2 ELISA’s for quantification of IgG1k, IgG1λ, IgG2k and IgG2λ, using recombinant IgG molecules as standards. The light-chain subclass ELISA were validated against the nephelometry data, which correlated well with the sum obtained from the ELISA as determined by regression analysis (Figure 4). Thus, nephelometry data fitted very well for IgG2 quantitatively (slope 1.10), but for IgG1 the nephelometry data seemed to be somewhat underestimated (slope 0.45), or overestimated by the ELISA (or both). In support of this, the average k/λ ratios were 1.4 ± 0.4 (range 0.7–2.3) and 2.4 ± 1.0 (range 0.8–4.5) for IgG1 and IgG2, respectively, suggesting IgG1λ values to be overestimated by the ELISA as total IgG k/λ ratios have been estimated to be around 2.0, which has never been determined for the IgG subclasses to our knowledge [27]. Importantly, the sum of the light chain ELISA values for either IgG1 or IgG2 k and λ correlated equally well with nephelometry data (Figure 4, R² = 0.73 and 0.75 for IgG1 and IgG2 respectively, p < 0.0001). Thus, we concluded that the inconsistency seen between the estimated levels for IgG1 by nephelometry and IgG1k+IgG1λ was constant over the whole concentration range, suggesting the possible skewing to be relative, and not affecting the relative IgG1k and IgG1λ ratios measured at either high or low IgG1 levels.

Using this method, we estimated that the transport of both IgG1k and IgG1λ was comparable with a cord/mother ratio of 1.6 (Fig. 4A–B). Placental transport of IgG2 was also equally efficient for both light chain variants (average IgG2 cord/mother ratio = 0.99 and 0.93 for κ and λ, respectively) (Fig. 5C–D). Again, the relative transport for the light chain isotypes (Fig. 5E) was comparable as it was for their respective IgG subclasses (Fig. 1A). There was a strong correlation between IgG1k and IgG1λ transport (R² = 0.860, p < 0.0001), and a weaker one between IgG1k and the IgG2 two isotypes (R² = 0.549, p < 0.0001 and R² = 0.698, p < 0.0001 for κ and λ, respectively) (Fig. 5E). Again, the relative transport for the light chain isotypes (Fig. 5E) was similar as observed for the IgG subclasses (Fig. 1A). The more IgG1k transported, the more IgG1λ was transported as well, which was also similarly increased, but to a lower level, for both IgG2 light chain isotypes as IgG1k transport correlated significantly with that of and IgG1λ (R² = 0.860, p < 0.0001), IgG2k (R² = 0.549, p < 0.0001), and IgG2λ (R² = 0.698, p < 0.0001)
No difference was found between the slopes for IgG2\textsubscript{k} and IgG2\textsubscript{l}. Although the slope for IgG1\textsubscript{l} differed significantly from IgG1\textsubscript{k} (P = 0.030), the difference was minimal, with slightly lower IgG1\textsubscript{l} transport at high IgG1\textsubscript{k} and vice versa (Fig. 5E). However, paired comparison between either IgG1\textsubscript{k} and IgG1\textsubscript{l} or IgG2\textsubscript{k} and IgG2\textsubscript{l} subclass yielded no significant difference. Also, the relative transport of IgG1\textsubscript{k} and IgG1\textsubscript{l}, as well as that for IgG2\textsubscript{k} and IgG2\textsubscript{l}, were not significantly different.

The half-life of IgG2 light chain isoforms does not differ in mice

As a previous study reported a different half-life for IgG2\textsubscript{k} and IgG2\textsubscript{l}, our next step was to measure the serum half-life of IgG2\textsubscript{k} and IgG2\textsubscript{l} in Balb/C mice injected with 200 mg of recombinant fully human IgG2\textsubscript{k} or IgG2\textsubscript{l} [15]. Concentration of human IgG in their serum was quantified at every 5 days over a two week period post injection. No difference was found between the clearance rates of IgG2\textsubscript{k} and IgG2\textsubscript{l} (Fig. 6A). We then tested if the clearance rate of IgG2\textsubscript{k} might be affected by the A/B isoforms recently described [4,6,28]. The different isoforms were deliberately generated as described previously (Fig. 6B) [5]. The clearance of neither IgG2\textsubscript{k} isoform differed from IgG2\textsubscript{l}, nor did the clearance of IgG2\textsubscript{KA} differ from IgG2\textsubscript{KB} in Balb/C mice (Fig. 6C). In addition, IgG2\textsubscript{KA} and IgG2\textsubscript{KB} were also cleared at equal rates in C57Bl/6 mice (Fig. 6D). C57Bl/6 FcRn KO mice cleared human IgG much faster than both WT expressing mice and displayed no difference between the two IgG2\textsubscript{k} isoforms (Fig. 6D). Antibody half-life was determined to be around 10 days in C57Bl/6 mice and around 4 days in Balb/C mice. In the FcRn KO mice, the levels had dropped below the detection limit of the assay at day 10. In conclusion, neither the half-life nor the transplacental transport is affected by the light chain isotype of IgG2 and thus not on structural isomerization of IgG2.

IgG2 half-life is identical between the light chain isotypes in humans

As these results are at odds with what we expected based on previously published work on a different, but mechanistically similar FcRn-mediated function, describing differences in half-life of IgG2\textsubscript{k} and IgG2\textsubscript{l} in Balb/C mice, we tested if the half-life of our IgG2 antibodies differed in humans [15]. We analyzed sera from IVIg-treated X-linked agammaglobulinemic patients as we described before, comparing the relative amounts found in the IVIg preparation itself to amounts found in patient serum four weeks later [20]. The relative amount remaining after IVIg-treatment was identical for IgG1\textsubscript{k} and IgG1\textsubscript{l}, but also for IgG2\textsubscript{k} and IgG2\textsubscript{l}, indicating that serum half-life does not differ between the light-chain isotypes in humans (Fig. 7B). Curiously, the relative amount remaining after 4 weeks was significantly increased for IgG2 in the IVIg-treated agammaglobulinemic patients (Fig. 7B), while it was significantly decreased for placental transport of IgG2 (Fig. 7A), indicating a fundamental difference between these two processes.

Figure 3. Determination of the FcRn binding properties of IgG1 and IgG2 light chain variants. Binding of titrated amounts of soluble human FcRn (62.5–8000 nM) at pH 6.0 to human IgG variants immobilized onto CM5 sensor flow cells. The relative affinity constants derived (KD) are indicated.
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FcRn mediates IgG transcytosis and recycling, both of which are compromised for IgG3, with reduced serum persistence and placental transport. We recently reported that these effects are attributable to a single amino acid difference between IgG3 and the other subclasses. While IgG1, IgG2 and IgG4 have a histidine at position 435, a key amino acid responsible for the pH-dependent binding to FcRn, most IgG3 allotypes have an arginine at this position, causing them to lose competition for recycling and transcytosis \[18,20\]. IgG3 allotypes which express a histidine at this position have equal half-life to that of IgG1, and are transported equally well across the placenta \[18,20\].

For IgG2, however, these FcRn-mediated transcytosis and recycling functions seem to diverge \[18,29\]. Although IgG1, IgG2 and IgG4 share all known contact residues with FcRn, and FcRn is known to mediate both IgG transcytosis and extend IgG serum persistence in a very similar manner, IgG2 crosses the human placenta with a markedly lower efficiency, with trans-placental transport more closely matching that of IgG3.

Interestingly, Montano and Morrison reported accelerated clearance for IgG2_\(\kappa\), compared to IgG2_\(\lambda\) in wild-type mice, indicating that the light chain may affect the interaction of IgG2 with FcRn \[15\]. This possibility became more plausible after the discovery that IgG2_\(\kappa\) uniquely exists in three structurally distinct isoforms, allowing for the possibility that a single isoform of IgG2_\(\kappa\)...

Figure 4. Validation of IgG1- and IgG2- light chain specific ELISA. Results from IgG1 total (A) and IgG2 total (B) were plotted against the sum of IgG1_\(\kappa\) and IgG1_\(\lambda\), or IgG2_\(\kappa\) and IgG2_\(\lambda\), respectively. The results of regression analysis are indicated in each panel, along with Pearson’s correlation. doi:10.1371/journal.pone.0108319.g004

Figure 5. Equal placental transport of \(\kappa\) and \(\lambda\) of IgG1 and IgG2 subclasses. IgG1_\(\kappa\) (A), IgG1_\(\lambda\) (B) and IgG2_\(\kappa\) (C) IgG2_\(\lambda\) (D) light chain isotype from sera in Fig. 1 were quantified by subclass- and light chain specific ELISA and each mother-child pair was plotted on the \(x\)- and

A) 

B) 

C) 

D) 

E)
may be influencing the overall interaction of IgG2 with FcRn.

We first investigated whether the difference between IgG2\(k\) and IgG2\(l\) could be found in paired mother-cord samples. Although the mother/cord ratio was significantly lower for IgG2 than for IgG1 or IgG4, there was no evidence that this was affected by the light chain isotype.

While the original finding that the half-life of IgG2\(k\) and IgG2\(l\) diverge, was observed for mouse-human chimeric antibodies in BALB/c mice, we investigated whether the same applied to fully human recombinant IgG2 antibodies, derived from an human IgG2\(l\) hybridoma but expressed with either \(k\) or \(l\) light chains, and tested them in both C57Bl/6 and BALB-c mice [30,31]. No differences were found neither in BALB/c nor C57Bl/6 mice. Curiously, we found differences in half-lives between experiments. In Balb/C mice we first found on average a half-life of approximately 6.5 days (Fig. 6A), while in another set of experiment it was down to 4.5 days (Fig. 6C). Furthermore, clearance seemed to be accelerated at later time points, as in Fig. 6A. In C57Bl/6 mice, the calculated half-life was again about 6.5 days. One possible reason is that FcRn expression (and consequently, IgG half-life) differs between different mouse strains, but perhaps an even more attractive explanation is that FcRn expression has been reported to be sensitive to NF-\(\kappa\)B signalling, suggesting FcRn-levels may also be under the control of the animal’s microbial status [32,33]. This might also explain

![Figure 6](https://example.com/figure6.png)

**Figure 6. No difference in half-life of IgG2 light chain isotypes \(k\) and \(l\) in mice.** (A) Recombinant human IgG2\(k\) and \(l\) in Balb/C mice was injected and measured by total IgG ELISA for a two week period following injection of 200 \(\mu\)g IgG. Calculated half-lives were \(7.2 \pm 1.48\) and \(6.4 \pm 0.84\) days for IgG2\(k\) and IgG2\(l\). (B) Enrichment of IgG2\(k\) isoforms was performed as described in Dillon et al 2008. HPLC elution profiles of IgG2\(k\)A and \(k\)B structural isomeres on a Dionex ProPac WCX-10 (4.0/250 mm) column are depicted. IgG2\(k\)B isoform was generated by incubation of 3 mg/ml IgG2\(k\) in 200 mM Tris buffer at pH 8 with 6 and 1 mM of cysteine and cystamine, respectively. For IgG2\(k\)A synthesis 0.9M guanidine hydrochloride (GuHCl) was also added. The samples were kept in the dark and placed at 4°C for 48–72 h. Following incubation the antibody was run on a Zeba spin desalting column (Pierce) for buffer exchange into PBS. (C) The clearance of IgG2\(k\), IgG2\(k\)A, and IgG2\(k\)B in Balb/C mice. Calculated half-lives were \(4.0 \pm 0.58\), \(5.39 \pm 0.85\), and \(3.7 \pm 1.04\) days for IgG2\(k\)A, IgG2\(k\)B and IgG2\(k\), respectively. (D) Clearance of IgG2\(k\)A and IgG2\(k\)B in WT and FcR\(n\) \(-/\) mice. Graphs in (A, C–D) depict mean and standard deviations of results obtained for 4 mice per group. Half-lives were calculated assuming exponential decay and reported in days ± standard error of means. No significant difference in half-life was detected between the two isotypes.

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Figure 7. Equal placental transport and serum clearance of IgG1 and 2 light chain isotypes κ and λ. (A) The average IgG1 and IgG2 placental transport (maternal/child) ratios were compared according to their light chain isotype. (B) Clearance of IgG1 and IgG2 κ and λ were quantified in serum by subclass- and light chain specific ELISA and subclass composition was compared to that found in the IVig used. No preferential clearance of one light chain isotype was detectable in either IgG subclass. 

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discerns between the two light chain isomers during intracellular transport.

This leaves the dilemma of explaining the lack of IgG2 transport across the placenta, observed in the paired mother-cord samples, compared to equal serum persistence. It has been postulated and subsequently disproved that FcγRIIb found on endothelial cells is involved in IgG transport across the murine yolk sac [34–38]. However, Mohanty and co-workers did not fully exclude the possibility that FcγRIIb may play a role in transplacental transport in humans [34–37]. If FcγRIIb is indeed involved, it may provide an explanation to the low transport of IgG2, as this is the only subclass with almost no measurable binding affinity to FcγRIIb [39,40]. However, the group of Ravetch has reported that sialylation of the N-linked glycan at position 297 in the IgG Fc affects binding to mouse FcγRIIb, resulting in 10× lowered binding affinity for both mouse IgG1 and IgG2a [41]. If FcγRIIb is indeed involved in placental transport, this would suggest a skewed transport of sialylated IgG across the placenta. However, we have found no such difference for any of the human subclasses, as all glycoforms of all subclasses were transported equally well across the placenta, suggesting FeRN, which does not require the Fc-glycans for binding, is sufficient for transplacental-IgG transport [42,43].

While it is conceivable that the requirements for FeRN-binding differ for IgG2 compared to the other subclasses due to the shorter hinge of IgG2 and the closer proximity of the Fab fragments to the Fc, there is no clear reason why this should affect FeRN-mediated recycling and transcytosis differently. The possibility exists that this extracellular binding also affects FeRN-mediated signaling, as transcytosis and recycling are regulated by different proteins, with actin motor myosin Vb and the GTPase Rab25 initiating transcytosis from the recycling endosome, while Rab11a regulates recycling to the basolateral membrane [44,45]. The role for the tryptophan-based basolateral-targeting signal identified in the FeRN cytoplasmic tail is also unknown [46]. In support of this view, we did observe that while IgG2 transport through the placenta is indeed low, its recycling and half-life extension in the human circulation is even better than for IgG1. If, and then how, these intracellular mechanisms are differently affected by the stoichiometry of the IgG2 molecules still needs to be elucidated.

In conclusion, we report and confirm that human IgG2 is not efficiently transported across the human placenta, despite a normal half-life. Although a previous report hinted at differential recycling of IgG2 depending on the light chain isotype, we found no preference for either recycling or placental transport of either IgG2κ or IgG2λ. In addition, the half-life extension in mice was not affected by the different IgG2κ structural isoforms. This is suggestive of alternative mechanisms, either involving an alternative receptor or mechanism differentiating between IgG transcytosis and recycling.

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Author Contributions

Conceived and designed the experiments: GV CES TR HKE JTA. Performed the experiments: HKE NMS TR JTA. Analyzed the data: HKE NME TR GV CES SS JTA. Contributed reagents/materials/analysis tools: HKE GV SS TR JTA. Wrote the paper: HKE NMS GV. Handled patients: SS. Obtained human samples for the study: SS. Read and critically edited the manuscript: HKE NME TR GV CES SS JTA.
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