Combined glyco- and protein-Fc engineering simultaneously enhance cytotoxicity and half-life of a therapeutic antibody

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Abbreviations: ADCC, antibody-dependent cell-mediated cytotoxicity; ADCP, antibody-dependent cell-mediated phagocytosis; FcRn, neonatal Fc receptor; FcγR, Fc gamma receptor; IgG, immunoglobulin G; mAb, monoclonal antibody; MS, cycle of mutagenesis and selection; NK, natural killer; PK, pharmacokinetics; RT, room temperature; SPR, surface plasmon resonance

While glyco-engineered monoclonal antibodies (mAbs) with improved antibody-dependent cell-mediated cytotoxicity (ADCC) are reaching the market, extensive efforts have also been made to improve their pharmacokinetic properties to generate biologically superior molecules. Most therapeutic mAbs are human or humanized IgG molecules whose half-life is dependent on the neonatal Fc receptor FcRn. FcRn reduces IgG catabolism by binding to the Fc domain of endocytosed IgGs in acidic lysosomal compartments, allowing them to be recycled into the blood. Fc-engineered mAbs with increased FcRn affinity resulted in longer in vivo half-life in animal models, but also in healthy humans. These Fc-engineered mAbs were obtained by alanine scanning, directed mutagenesis or in silico approach of the FcRn binding site. In our approach, we applied a random mutagenesis technology (MutaGen™) to generate mutations evenly distributed over the whole Fc sequence of human IgG1. IgG variants with improved FcRn-binding were then isolated from these Fc-libraries using a pH-dependent phage display selection process. Two successive rounds of mutagenesis and selection were performed to identify several mutations that dramatically improve FcRn binding. Notably, many of these mutations were unpredictable by rational design as they were located distantly from the FcRn binding site, validating our random molecular approach. When produced on the EMABling™ platform allowing effector function increase, our IgG variants retained both higher ADCC and higher FcRn binding. Moreover, these IgG variants exhibited longer half-life in human FcRn transgenic mice. These results clearly demonstrate that glyco-engineering to improve cytotoxicity and protein-engineering to increase half-life can be combined to further optimize therapeutic mAbs.

Introduction

Therapeutic monoclonal antibodies (mAbs) of the immunoglobulin G (IgG) isotype are in widespread use for the treatment of a variety of disorders, including cancers, autoimmune and infectious diseases. Since the approval of the first-generation mAbs in the late 1990s, tremendous improvement of their efficacy has been achieved by molecular engineering. The second-generation of therapeutic mAbs usually included optimized variable domains (Fv) such as human/humanized Fv domains or higher antigen affinity. Over the past decade, a third-generation of mAbs has emerged1 with improved Fc region because it was clearly established that the success of mAbs as cancer therapeutics substantially relied on their ability to engage the immune system via this Fc domain.2,3 This Fc-dependent engagement is mediated by activation of the lytic complement pathway (classical pathway) and crosslinking of Fcγ receptors (FcγRs) widely expressed on many innate immune effector cells such as natural killer (NK) cells, monocytes, macrophages and neutrophils.4 Therefore, current knowledge of the mechanisms of action of therapeutic mAbs involve direct (blockade of intracellular signaling, deprivation of growth hormone or pro-apoptotic effect) as well as indirect (complement- or cell-mediated cytotoxicity) killing of the tumor cells via Fv and Fc domains respectively.5 With the exception of T cells, immune effector cells express one or more activating FcγRs (FcγRI, FcγRIIA, and FcγRIIIA) and, for the majority, co-express the inhibitory FcγR (FcγRIIB). Following engagement of their activating FcγRs, these cells kill tumor cells through antibody-dependent cell-mediated cytotoxicity (ADCC) or through phagocytosis (ADCP).

The rationale that improved mAb-mediated effector functions will be clinically beneficial is supported by the...
correlation of clinical outcomes of mAb therapies in patients with natural polymorphisms of FcγRs. The allelic FcγR variants show different affinities for the Fc domain of IgG1 and more favorable clinical responses were observed in patients who were homozygous for the higher-affinity allele of the activating receptor FcγRIIIA (V158). This has been observed for treatment with the anti-CD20 rituximab in follicular lymphoma, the anti-HER2 trastuzumab in metastatic breast cancer and the anti-EGFR cetuximab in metastatic colorectal cancer. These findings, obtained for various specificities and malignancies, suggest that the low-affinity receptor FcγRIIIA is crucial for the anti-tumor activity of mAbs in vivo and that increasing crosslinking of this receptor should translate into enhanced therapeutic activity. To this aim, glyco-engineered mAbs have been designed. Indeed, the absence of α1,6 innermost fucose residues on the Fc glycan moiety of mAbs has long been associated with enhanced affinity for FcγRIIIA, and there is strong evidence that such afucosylated or low fucosylated mAbs translate into enhanced therapeutic activity in mouse models in vivo. To date, more than 20 third-generation glyco-engineered mAbs, with higher ADCC, are being evaluated in clinical studies. Two of these mAbs have already been approved, mogamulizumab in March 2012 and obinutuzumab in November 2013, confirming the success of this approach.

In addition to increasing mAbs cytotoxicity, extensive efforts have also been made to improve mAbs pharmacokinetic (PK) properties. Despite the relatively long half-lives of mAbs, improved molecules with longer half-lives and conserved efficacy would permit less frequent dosing schedules and provide higher patient convenience. IgG half-life is dependent on the neonatal Fc receptor (FcRn), a member of the MHC class I superfamily, which is expressed by many cell types including endothelial cells, macrophages, monocytes and monocyte-derived dendritic cells in human. FcRn is a high affinity Fc receptor that binds IgG in a strictly pH-dependent manner. IgGs are internalized by non-specific pinocytosis into the endosomes of cells where the low pH (pH 5–6) promotes binding to FcRn with nanomolar affinity. Bound IgG-FcRn complexes are recycled back to the cell surface and dissociate at the neutral pH of the extracellular fluid, returning to the blood circulation. IgGs that do not bind to FcRn traffic into the lysosomes where they are degraded by proteases. Moreover, FcRn was shown to be involved in IgG transport across the feto-maternal barrier and across endothelial and epithelial cellular barriers increasing their overall bioavailability.

This property has been successfully exploited for pulmonary delivery, as well as intranasal immunization with Fc-fusion proteins. Lastly, FcRn was also reported to transport IgG-antigen complexes, thus favoring antigen-presentation, and to enable phagocytosis of IgG-opsonized bacteria by neutrophils, demonstrating a wide range of functions for this receptor in several immunological and non-immunological mechanisms.

Consistent with its role in IgG catabolism, several Fc-engineered mAbs with increased FcRn affinity and conserved pH dependency were designed and resulted in longer half-life in vivo in human FcRn transgenic mice (hFcRn mice) and cynomolgus monkeys. Hence, Fc-engineered variants (Fc-LS) of the anti-VEGF bevacizumab and the anti-EGFR cetuximab demonstrated half-lives extended up to almost 5-fold in hFcRn mice and 3-fold in cynomolgus monkeys. These mAbs exhibited greater anti-tumor activity when tested in hFcRn/Rag1−/− tumor xenografted mice, validating that prolonged exposure due to FcRn-mediated enhancement of half-life should also improve therapeutic mAbs efficacy in human. Moreover, for the first time in healthy humans an Fc-engineered anti-RSV (motavizumab-YTE) recently demonstrated a significant increase in serum half-life of up to 100 days, validating the usefulness of increasing FcRn binding and fully confirming the results previously obtained in animal models.

For the past 20 years, the Fc/FcRn interaction has been extensively studied by directed mutagenesis and crystallography using the respective human, murine and rat proteins. The FcRn binding site is structurally conserved among species and comprises three distinct zones in the CH2 and CH3 domains that are localized in the CH2/CH3 interdomain region. Fc-engineered mAbs with increased FcRn affinity were obtained by alanine scanning, but also directed mutagenesis or in silico approaches focused on the FcRn binding site. Therefore, the resulting Fc variants usually include 2–3 substitutions in or close to the FcRn binding site. To identify new mutations that could be distant from the FcRn binding site but have an indirect, and possibly cooperative, positive impact on the Fc/FcRn complex, we randomly mutated the entire Fc domain and selected several FcRn affinity enhanced Fc variants at pH 6.0. To improve both mAb cytotoxicity and half-life, we produced the selected IgG variants as low fucosylated molecules and demonstrated increased serum persistence in hFcRn mice, as well as conserved enhanced ADCC.

Results

First round of mutagenesis and selection (MS1)

Construction of random libraries

The human IgG1 Fc gene encoding amino acid residues 226–447 (EU index as in Kabat), referred to as Fc wild type (Fc-WT), was cloned into our modified phagemid vector. This Fc fragment includes the entire CH2 and CH3 domains of the last 5 amino acids of the hinge region. To set up the experiments, the double variant T250Q/M428L (Fc-QL) and the triple mutant M252Y/S254T/T256E (Fc-YTE) were constructed as positive controls. In the phagemid vector, several fully randomized libraries were generated using the MutaGen™ technology that uses low fidelity human DNA polymerases (mutases) to introduce random mutations homogeneously throughout the target sequence. Three distinct human mutases (pol. β, pol. η and pol. ι) were used in different conditions to create complementary mutational patterns. Four different sub-libraries were constructed for this first round of mutagenesis and selection (MS1). A first sub-library was obtained using pol. β on the Fc-WT gene and contained 3.2 × 10^6 clones (called Mut1.1). The DNA of this first sub-library was used to generate the second and the third sub-libraries, using respectively pol. β (3.8 × 10^6 clones, Mut1.2) and pol. η plus ι (3.0 × 10^6 clones, Mut1.3). The fourth library was generated with pol. η alone
on the Fc-WT gene ($1.0 \times 10^6$ clones, Mut1.4). These four sub-libraries were proportionally mixed to obtain the final library Mut1, representing $1.1 \times 10^7$ clones (Table 1).

For each sub-library and the final pooled library Mut1, 48 to 192 clones were sequenced after PCR on cells and sequences were analyzed using MilleGen’s proprietary software Mutanalyse4Fc adapted for the Fc molecule from the Mutanalyse 2.5 software described previously. Analysis of the four sub-libraries’ sequences showed that the use of two sequential mutagenesis steps (Fc-WT to Mut1.1 and Mut1.1 to either Mut1.2 or Mut1.3) permitted an increase in the mutation rate (Table 1). The mutation rate of Mut1.1 is 4.95 nucleotide mutations per kilo base (kb), whereas Mut1.2 and Mut1.3 had mutation rates of 8.04 and 8.16 mutations per kb, respectively. The percentage of mutated sequences was also increased from Mut1.1 (66.7%) to the two sub-libraries (96.4 and 81.7%, respectively). The mutation rate for Mut1.4, constructed with pol. η alone, is high (13.89 mutations per kb) but this library contains only 40% of mutated sequences. For Mut1.3, 22 sequences for Mut1.4 and 35 sequences for Mut1) showed that the frequency of nucleotide mutations of Mut1 library is 7.31 mutations per kb, which corresponds to 4.89 mutations per Fc gene [666 nucleotides (nt)] (Table 1). Among these mutations, 84.0% are substitutions, 14.2% are deletions and 1.8% are additions, these last two categories introducing frame shifts in the gene. When considering only the sequences mutated in frame, which represent 42.0% of the clones (excluding the unmutated sequences and the sequences with a stop codon other than a TAG), the mutation frequency is 4.05 mutations per kb, corresponding to 2.63 mutations per gene (1 to 6 mutated nucleotides per gene), with mainly 1 mutation per gene (Fig. 1A). The mutation analysis was also performed at the protein level. The Mut1 library contains 27.5% of clones expressing the Fc-WT [unmutated or with silent mutation(s)], 37.3% of clones containing a sequence out of frame or with a stop codon (different from the amber stop codon TAG expressed in our system) and 35.1% of clones with a mutated sequence (Fc variants). These last clones represent the active part of the library, comprising $-3.8 \times 10^6$ mutated clones with on average 2.35 mutated amino acids per molecule. The distribution of the mutations on the Fc sequence was also analyzed and confirmed that the random mutations are evenly distributed along the entire gene without any hot spot (Fig. 1B).

![Figure 1](https://example.com/figure1.png)

**Figure 1.** Analysis of the mutated sequences in frame in the Mut1 library. (A) Distribution of the number of nucleotide mutations per Fc. (B) Distribution of the mutations along the Fc region (amino acids).

| Library name | Mut1.1 | Mut1.2 | Mut1.3 | Mut1.4 | Mut1 | Mut2 |
|--------------|--------|--------|--------|--------|------|------|
| Template     | Fc-WT  | Mut1.1 | Mut1.1 | Fc-WT  | Fc-WT or Mut1.1 | 42 variants |
| Mutase       | pol. η | pol. η | pol. η | pol. η | pol. η, pol. η, pol. η | pol. η, pol. η |
| Diversity (clones) | $3.2 \times 10^6$ | $3.8 \times 10^6$ | $3.0 \times 10^6$ | $1.0 \times 10^6$ | $1.1 \times 10^7$ | $2.0 \times 10^7$ |
| Number of analysed sequences | 120 | 83 | 82 | 130 | 276 | 86 |
| Mutated sequences (%) | 66.7 | 96.4 | 81.7 | 40.0 | 78.3 | 83.7 |
| Mutation rate* (nt/kb) | 4.95 | 8.04 | 8.16 | 13.89 | 7.31 | 4.54 |
| Substitutions (%) | 82.9 | 80.1 | 85.2 | 86.1 | 84.0 | 96.3 |
| Deletions (%) | 15.2 | 17.5 | 12.1 | 11.7 | 14.2 | 3.2 |
| Insertions (%) | 1.9 | 2.4 | 2.7 | 2.2 | 1.8 | 0.5 |

*excluding the unmutated sequences

![Table 1](https://example.com/table1.png)
Selection of Fc variants on human FcRn and characterization by phage-ELISA

The Fc variants of the Mut1 library were displayed on the M13 bacteriophage using the classical technique of phage display. Selection was performed using three different strategies, two using immunoplates coated with either biotinylated FcRn or the FcRn-p3 fusion molecule and one using biotinylated FcRn in solution trapped with streptavidin-coated magnetic bead, to increase the chance to select for affinity-enhanced FcRn binders. FcRn binding and washing steps were performed at pH 6.0 whereas bound Fc-phages were eluted at pH 7.4 to preserve Fc/FcRn interaction pH-dependency, rendering the selection process highly specific as well. Three to eight rounds of selection were performed for each strategy. Retrieved clones were isolated, sequenced and analyzed with internally developed bioinformatics tools.

During the MS1 process, a total of 227 different Fc variants were isolated. These 227 mutated Fc-phages were then ranked for their FcRn binding using a comparative phage-ELISA assay at pH 6.0 on FcRn-coated plates with the Fc-WT as reference. In this phage-ELISA assay, the Fc-QL and Fc-YTE positive controls, expressed on the M13 bacteriophage, showed a better binding than Fc-WT and similar pH-dependent binding properties, i.e., strong binding at pH 6.0, decreased binding at higher pH and no binding at pH 7.4, validating our phage display system (data not shown). Fc-QL and Fc-YTE had a specific signal that was on average 3-fold stronger than the Fc-WT (ratio Fc-QL/Fc-WT = 3.2, Fc-YTE/Fc-WT = 3.5). From the 227 mutants isolated during MS1, 139 were considered positive with a ratio/Fc-WT > 2 and 73 were better than the positive control Fc-QL (ratio/Fc-WT > 3.2). The best clone, S5A-41, had a ratio/Fc-WT of 9.0 (Fig. 2A). Sequence analyses of these positive Fc variants showed that they contain on average 2.7 mutated amino acids per Fc and permitted to identify 4 key positions, which are mutated in more than 10% of the positive clones. (C) Sequences of the 10 best clones isolated during MS1. Key positions as identified in Figure 2B are in bold.

Second round of mutagenesis and selection (MS2)

Construction of random libraries

Based on the results obtained during MS1, we performed a second round of mutagenesis and selection (MS2) to further improve the Fc/FcRn binding. As previously described for MS1, randomized libraries were constructed with the MutaGen™ technology. For these new libraries, the DNA template used was a DNA pool of 42 variants isolated during MS1 and having improved FcRn binding by phage-ELISA. To minimize the number of mutations of the final variants, only the variants with one or two amino acid mutations were included in this second mutagenesis process. In addition, variants mutated on position N434 were also excluded as the mutations N434Y or S were too predominant and not novel, our aim being to identify new mutations. At the protein level, our pool of 42 single and double
variants has a mutation rate of 1.45 amino acids per Fc. As many variants have silent mutations (15 variants with one silent mutation, 4 variants with 2 and 4 variants with 3 silent mutations), the nucleotide mutation rate of the pool is on average of 3.43 mutations per kb, i.e., 2.28 mutations per gene (1 to 5 nucleotide mutations, Fig. 3B). Two libraries were constructed with mild conditions to achieve a low mutation rate. A first library was obtained using pol. β (1.9 x 10^7 clones, Mut2.1) and a second library with pol. η (1 x 10^6 clones, Mut2.2). These two libraries were proportionally mixed to obtain the final library Mut2, representing 2 x 10^7 different clones (Table 1). The frequency of mutations of the Mut2 library is 4.54 nucleotide mutations per kb, i.e., 3.02 mutations per gene (666 nt). Among these mutations, 96.3% are substitutions, 3.2% are deletions and 0.5% are additions (Table 1). When considering only the sequences in frame, the mutation frequency increases to 4.34 mutations per kb, i.e., 2.89 mutations per gene (1 to 7 mutated nucleotides per gene, Fig. 3B). At the protein level, the Mut2 library contains 17.4% of clones expressing the Fc-WT fragment (unmutated or with silent mutations), 9.3% of clones with a sequence out of frame or with a stop codon (not expressed) and 73.3% of clones with a mutated sequence (Fc variants). These last clones represent the active part of the library comprising 1.5 x 10^7 different clones with an average of 1.87 mutated amino acids per molecule.

Selection of Fc variants on human FcRn and characterization by phage-ELISA

As previously described for MS1, the Fc variants of Mut2 library were displayed on M13 bacteriophages and were selected on human recombinant FcRn using the three different strategies described above (solid and liquid phases). During the MS2 process, 223 different mutated clones were isolated and ranked by phage-ELISA on FcRn coated plates at pH 6.0. Because the difference between the signals of the Fc-WT and the Fc variants was too great to be compared on the same ELISA plate, the Fc variants were directly compared with the Fc-QL variant as reference. Moreover, the best Fc variant isolated during MS1, S5A-41, was used as positive control on each phage-ELISA plate. Among the 223 Fc variants tested, 209 Fc variants were better than the Fc-QL (ratio/Fc-QL at least > 1.1) and 39 Fc variants were better than the S5A-41 variant. The best clone isolated during MS2, C6A-69, had a ratio/Fc-QL of 9.0 (Fig. 4A). These results emphasize the importance of performing this second round of random mutagenesis to further improve the binding. To compare the variants with the Fc-WT, an estimated ratio/Fc-WT was calculated by multiplying the ratio/Fc-QL of the variants with the ratio/Fc-WT of the Fc-QL (ratio/Fc-WT = 3.2 x ratio/Fc-QL) determined during MS1. Overall, 219 Fc variants were considered positive with an estimated ratio greater than 2 compared with the Fc-WT. Sequence analysis of these positive Fc variants showed that they contain on average 3.9 mutated amino acids per Fc and permitted identification of 9 key positions (P230, F241, V264, T307, N315, A330, A378, N389, and N434), including the 4 positions previously identified and 5 new positions (Fig. 4B). One position is in the FcRn binding site and is well-characterized (T307), the other positions are in the CH2 and CH3 domains but not directly in the FcRn binding site. Three positions are nonetheless close to the FcRn binding site (N315, A378 and N389). Finally, we found the position N434 mutated in the majority of the positive variants, as exemplified for the 10 best Fc variants isolated (Fig. 4C).

Production and functional characterization of IgG variants

Choice and production of IgG variants

To validate the results obtained by phage display, we first produced Fc variants as Fc fragments in E. coli. For that purpose, we chose 15 Fc variants with distinct enhanced FcRn binding properties as shown by phage-ELISA. Overall, FcRn binding analysis of Fc fragments by ELISA and SPR confirmed the phage-ELISA ranking and pH-dependent binding of the selected Fc variants (data not shown). We then chose six among the ten best Fc variants selected by phage display (Fig. 4C; Table 2). These Fc variants correspond to the 5 best variants plus the C6A-66 variant, which has the originality of containing an amino acid deletion (named E294Del) that we were keen on characterizing. Given the relatively high number of mutations of these Fc variants, we then tried to ensure that each mutation had a real positive effect on the FcRn binding. To that aim, we constructed a database to store and analyze all the DNA sequences and phage-ELISA results obtained for the Fc variants isolated during MS1 and MS2 processes (~450 clones). Analysis of this database showed that many of the Fc variants characterized were combinations of 2–3 mutations also found in our 6 preferred variants. We therefore identified mutations that could be dispensable for improved FcRn binding. In an attempt to reduce the final number of mutations of the IgG variants, we constructed several new variants by removing these mutations and tested them by phage-ELISA as previously described. Most of the new variants showed a decreased FcRn binding compared with the starting variants. Finally, only one new variant was retained, C6A-78A, because the removal of one mutation from the C6A-78 variant (N383S)
seemed to have no effect on FcRn binding (data not shown). The C6A-78A variant, the 6 chosen variants (Table 2), the IgG-WT and the positive control IgG-YTE were produced as full-length IgGs with the variable regions of the chimeric anti-human CD20 CAT 13.6E12. To combine the advantages of glyco- and protein-Fc engineering, the IgG variants were produced in YB2/0 cells (EMABling® platform), permitting low fucosylation of the glycan moiety present in the mAb and thus enhanced ADCC as previously described for the anti-CD20 mAb ublituximab37-39 and the anti-RhD mAb roledumab.40,41 Before any subsequent biological test, the quality of the purified IgGs was assessed using several biochemical techniques ensuring that the biophysical behavior of the IgGs were not modified due to the mutations introduced (i.e., similar pI and molecular weights, aggregation rates < 5% and endotoxin levels < 7UI/mg).

### FcRn binding characterizations

The 6 IgG variants were compared with the IgG-WT for their FcRn binding properties using 3 different techniques: (1) ELISA using microtiter plates coated with FcRn, (2) binding to cells stably expressing a membrane version of FcRn, and (3) SPR on immobilized FcRn (Table 2; Fig. 5A and B). Whatever the technique used, all the IgG variants exhibited improved FcRn binding at pH6.0. By ELISA, the EC50 of IgG variants were at least 5.2-fold lower than that of the IgG-WT. The best results were obtained for the variants C6A-69 and T5A-74 (ratio EC50 variant/WT = 10.8 and 9.3, respectively). These data are in good correlation with the phage-ELISA results obtained previously to rank the variants isolated during MS1 and MS2, the C6A-69 variant being the best variant characterized during MS2. In the ELISA tests, the IgG variants displayed similar results, with

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**Table 2.** Comparison of binding of IgG variants to FcRn using 3 different techniques: ELISA on coated FcRn, FcRn binding on cells and SPR on immobilized FcRn

| IgG variants | Mutations | Phage-ELISA | ELISA on coated FcRn | FcRn binding on cells | Steady-state SPR measures on FcRn |
|-------------|-----------|-------------|----------------------|----------------------|-------------------------------|
| WT          | None      | Ratio OD variant/WT | EC50 (ng/ml) | Ratio EC50 WT/variant | IC50 (μg/ml) | Ratio IC50 WT/variant | KD (nM) | Ratio KD WT/variant |
| C6A-69      | T307A/N315D/A330V/E382V/N389T/N434Y | 1.0 | 11060.0 | 1.0 | 249.3 | 1.0 | 78.3 | 1.0 |
| C6A-78      | T256N/A378V/S383N/N434Y | 28.4 | 1021.6 | 10.8 | 1.4 | 178.1 | 16.8 | 2.8 |
| T5A-74      | N315D/A330V/N361D/A378V/N434Y | 27.8 | 1440.9 | 7.7 | 3.8 | 65.6 | 17.6 | 4.4 |
| T5A-70      | V259N/N315D/N434Y | 27.6 | 1191.8 | 9.3 | 1.4 | 178.1 | 10.6 | 7.4 |
| T5A-75      | P230S/N315D/N434Y | 27.2 | 2161.0 | 5.2 | 1.7 | 146.6 | 12.9 | 6.1 |
| C6A-66      | E294de/T307P/N434Y | 26.8 | 1904.0 | 5.8 | 1.5 | 166.2 | 13.8 | 5.7 |

EC50: Concentration at 50% saturation obtained from the ELISA binding curve of each IgG variant. The ratio is the EC50 of the IgG-WT divided by the EC50 of the variant. IC50: Concentration at 50% inhibition of the MFI signal obtained with Alexa-conjugated Rituximab alone (100%). The ratio is the IC50 of the IgG-WT divided by the IC50 of the variant.
ratios over IgG-WT ranging from 5.2 to 10.8, corresponding to a maximum ratio difference of 2.1. As expected, using the same protocol, no binding was detected at pH7.4 for any of the IgG variants (data not shown).

In a second series of tests, the IgG variants were assayed for their ability to bind FcRn expressed on cells. To this aim, we used Jurkat T cells stably expressing a truncated form of the human FcRn α-chain at their surface, similarly to what was recently described.42 This competitive assay clearly revealed the dramatic FcRn binding increase of the IgG variants compared with the IgG-WT (Fig. 5A). The measured IC50 for the IgG variants are significantly lower than that of the IgG-WT (Table 2). The IC50 decrease for the IgG variants compared with the IgG-WT ranged from 23.7 (for C6A-66) to 178.1 (C6A-67 and C6A-78). While similar results for all IgG variants were obtained in ELISA and SPR assays, this competitive cell-based assay revealed a different behavior for the C6A-66 deletion variant (factor 7.5 between C6A-69 and C6A-66). Currently, we have no explanation for this finding.

Lastly, human FcRn affinities of the IgG variants were measured by SPR at pH6.0. To rapidly compare several variants, we choose an experimental set up with recombinant human FcRn immobilized on the chip. As a result, IgG-FcRn affinities were higher than in the opposite format with immobilized mAbs, as previously described.43,44 In this format, the enhanced affinity of the IgG variants for human FcRn relative to the IgG-WT were predominantly driven by increased association kinetics as illustrated by the sensorgrams (Fig. 5B). In addition, as variable results were obtained when using the 1:2 model (included in the BIAevaluation software), we measured binding affinities at steady-state (K_D values). As illustrated in Table 2, the binding affinity of the IgG-WT for human FcRn was 78.3 nM and the K_D values of the 6 IgG variants ranged from 10.6 to 18.8 nM. Thus, in this test, the IgG variants exhibited similarly improved affinity, with increase ratio (WT K_D divided by variant K_D) between 4.2 and 7.4-fold. Finally, the positive control IgG-YTE, as well as the C6A-78 and C6A-78A variants, were tested in comparison with IgG-WT in a second complementary SPR experiment (Fig. 5C). This experiment confirmed that the C6A-78A variant has a similar affinity as C6A-78, albeit having one mutation less (N383S). Moreover, the IgG-YTE has a lower affinity increase than our IgG variants (ratio K_D WT/K_D YTE = 1.9), a result in line with the data obtained previously by phage-ELISA (Fig. 2A).

**Figure 5.** FcRn binding of the IgG variants. (A) FcRn binding of IgG variants on FcRn expressing cells. (B) Sensorgrams of FcRn binding at pH 6.0 for IgG-WT and C6A-66. Concentrations tested were 0, 2, 10, 25 (2 times), 50, 100, and 200 nM. (C) Complementary SPR measures for IgG-WT, IgG-YTE, C6A-78, and C6A-78A.
failed to bind hFcγRIIIA. Analogous results were obtained for hFcγRIIIA V158 allotype (data not shown). The IgG variants were also tested for their capacity to elicit ADCC with purified NK cells against CD20-expressing target cells (Fig. 6B). All IgG variants display similarly high ADCC, compared with RTX, except for the C6A-66 variant that has no ADCC activity. This variant contains three mutations, two substitutions directly located in the FcRn binding site (T307 and N434) and one deletion (E294 deleted), the latter most probably being responsible for the loss of hFcγRIIIA binding and ADCC activity. Overall, the ADCC results are in total agreement with the hFcγRIIIA binding capacity, showing altogether that five IgG variants have retained glyco-engineered improved ADCC activity.

Serum persistence in human FcRn transgenic mice

We then tested whether the improved FcRn binding properties of the IgG variants observed in vitro resulted in longer serum persistence in vivo. We therefore performed PK experiments in hFcRn mice that are homozygous for a knockout allele of murine FcRn and heterozygous for a human FcRn transgene (mFcRn-/- hFcRnTg 276 heterozygote on a B6 background). For these PK studies, each animal received a single intravenous injection of mAbs at 5 mg/kg via retro-orbital sinus, in a protocol similar to the one described previously. Blood samples were collected from the retro-orbital sinus at multiple time points and titrated by ELISA. The IgG variants were systematically compared with the IgG-WT and to IgG-YTE, a positive variant whose half-life has been largely studied in hFcRn mice earlier and was recently confirmed in humans.

Three different PK studies were performed (PK1 to PK3, parameters in Table 3). The IgG variants tested were those of Table 2 except for the C6A-78 variant replaced by the C6A-78A variant, which exhibited similar FcRn binding and ADCC properties but had one mutation less. In all three experiments, the IgG-WT exhibited a half-life of -1 d (23 h), a value lower than those published for other mAbs. This fast clearance in mice could be due to the high isoelectric point of the mAb (pI = 9.2), a parameter that has been shown to negatively affect half-life and clearance. In this set-up, all IgG variants displayed substantial serum half-life increase with ratios (half-life variant/WT) between 1.8 and 2.8. Similarly, IgG-YTE had a ratio between 2.3 and 3.3, slightly lower than the published values of -3.8. The IgG T5A-74 variant was tested in two different experiments (PK2 and PK3) and similar results were obtained (ratio 2.5 and 2.2, respectively). In the second experiment (PK2), the C6A-66 variant is slightly more persistent than IgG-YTE (ratio 2.8 vs. 2.3), whereas the C6A-74 variant demonstrates similar persistence as IgG-YTE (ratio of 2.3 for both).

Discussion

Given the remarkable role of FcRn in IgG serum persistence, IgG/FcRn interactions have long been thoroughly studied at the molecular level by crystallography and directed mutagenesis. The FcRn binding site in the Fc domain comprises key residues located at the CH2/CH3 domain interface: residues 252–254, 307 and 309–311 in the CH2 domain, and residues 433–436 in the CH3 domain. In addition, the pH dependence has been attributed to two histidines (H310 and H435) involved in salt bridges with acidic FcRn residues. Without the help of this molecular knowledge, we identified several mutations that have a positive impact on the IgG/FcRn complex using a fully random mutagenesis approach of the entire human Fc fragment combined with a robust selection process. Interestingly, most of the best variants isolated include one mutation in the FcRn binding site, N434Y or N434S, combined with original mutations that are close to or distant from the FcRn binding site. These mutations, mainly P230S/T, F241L, V264A/E, N315D, A330V, N361D and A378V/T, probably...
have a positive long-range effect on the overall structure of the Fc domain favoring the IgG/FcRn complex. Using molecular modeling, impact of these mutations was difficult to interpret so they would have been impossible to be predicted (data not shown). Furthermore, the effect of the combination of several mutations would be even more difficult to predict, as opposite, additive or synergistic effects can be obtained. All these results validate our engineering approach involving two successive rounds of random mutagenesis and selection.

On the other hand, the precise knowledge of the IgG/FcRn interactions have been successfully used before by others to design and isolate various Fc variants with improved FcRn affinity. Using molecular modeling, impact of these mutations was difficult to interpret so they would have been impossible to be predicted (data not shown). Furthermore, the effect of the combination of several mutations would be even more difficult to predict, as opposite, additive or synergistic effects can be obtained. All these results validate our engineering approach involving two successive rounds of random mutagenesis and selection.

On the other hand, the precise knowledge of the IgG/FcRn interactions have been successfully used before by others to design and isolate various Fc variants with improved FcRn affinity. For instance, residues 250 and 428 have been targeted for mutation due to their proximity to the CH2/CH3 interface and resulted in the isolation of the double variant T250Q/M428L (Fc-QL) with a ~27-fold binding increase on human FcRn expressed on cells. We successfully used this Fc-QL variant as a positive control for our phage display process. Nevertheless, contradictory results were obtained with this IgG-QL variant in PK studies in rhesus and cynomolgus monkeys. We therefore choose the IgG-YTE variant (M252Y/S254T/T256E) as a positive control for PK studies. This triple variant was obtained by phage display selection against murine FcRn of rationally designed libraries targeting Fc residues in or close to the FcRn binding site (residues 251–256, 308–314, 385–389, and 428–436). Introduced in a humanized IgG1, these mutations increased the binding to human FcRn by about 10-fold, resulting in a nearly 4-fold increase in serum half-life in cynomolgus monkeys. We therefore choose the IgG-YTE variant (M252Y/S254T/T256E) as a positive control for PK studies. This triple variant was obtained by phage display selection against murine FcRn of rationally designed libraries targeting Fc residues in or close to the FcRn binding site (residues 251–256, 308–314, 385–389, and 428–436). Introduced in a humanized IgG1, these mutations increased the binding to human FcRn by about 10-fold, resulting in a nearly 4-fold increase in serum half-life in cynomolgus monkeys. More importantly, when introduced into the anti-RSV motavizumab, the IgG-YTE variant resulted in a 2- to 4-fold longer half-life than the WT motavizumab in healthy humans. This is a crucial result as it confirms for the first time in humans the concept of half-life extension using Fc-engineered antibodies. On the other hand, in silico analysis was also used to design several variants combining mutations in or close to the FcRn binding site. Two variants, M428L/N434S (IgG-LS, Xtend) and V259I/V308F/M428L (IgG-IFL), showed respectively a 1.6- and 2.7-fold improved binding to human FcRn by SPR compared with the IgG-YTE. Nonetheless, similarly to what we observed in PK studies in FcRn mice, these variants exhibited only moderate half-life increase compared with the IgG-YTE, at best a 1.16-fold increase for the IgG-LS and a 1.30-fold increase for the IgG-IFL in one experiment. However, as we also observed, results can be variable since in a second experiment of this study no half-life increase was observed for the IgG-IFL compared with the IgG-YTE. Similarly, a lack of direct linear correlation between increased FcRn affinity and improved half-life had been observed earlier with the two variants N434A and T307A/E380A/N434A. These variants were isolated by alanine scanning mutagenesis and resulted in a 2.2-fold increase in hFcRn mice, whereas affinity differences were greater, respectively 3.4 and 11.8-fold increase compared with the IgG-WT. All these results seem to demonstrate that Fc engineering may have reached a plateau for FcRn improvement designed to increase half-life.

This in vivo limit could be due to the critical need to keep the pH dependency property of the IgG/FcRn interaction. Indeed, increased affinity at high pH could be potentially problematic by preventing the release of FcRn-bound IgGs. In accordance with this mechanism, augmented clearance was observed for IgG engineered for improved FcRn affinity at both pH 6.0 and pH 7.4, where pH dependence is reduced. Furthermore, recent studies clearly showed the difficulty of conserving the pH dependency because globally when a variant's affinity to FcRn is increased at pH 6.0, its binding at higher pH increases accordingly. Several mutations which resulted in a slight increase in FcRn binding at pH 7.4 were described, including M252Y, M252W, V308P, N434Y, and N434W. This last mutation, N434W, induces the highest FcRn affinity at pH 7.4 and despite an 80-fold increase binding to cynomolgus monkey FcRn at pH 6.0, serum half-life was similar to the IgG-WT. On the other hand, the mutation M252Y is included in the IgG-YTE variant but clearly resulted in
increased half-life for various mAbs.25 Likewise, our Fc variants include the mutation N434Y and herein we show that several variants have increased half-life in hFcRn mice. Combinations of several mutations may result in different pH dependency and very fine differences could have unpredictable effects on serum half-life. Therefore, pH dependency being difficult to assess and interpret we choose to directly test several variants for extended half-life in hFcRn mice. These FcRn-humanized mice, unlike wild-type mice, have been shown to be a reliable surrogate for studying human IgG serum half-life.23,53 Despite reduced circulating murine IgGs, these humanized mice reliably predicted antibody half-lives in primates when challenged with clinical mAbs for which PK data were available.24 Moreover, whereas human IgG binding and pH dependency are highly similar for human and monkey FcRn, human and murine FcRn have significant molecular differences rendering wild-type mice an inadequate model for studying half-life of engineered human IgGs.55,56 For instance, the N434A variant has increased serum half-life in monkeys, in correlation with its pH dependent increased binding to monkey FcRn, whereas in wild-type mice this variant has the same PK profile as the IgG-WT, due to unchanged binding properties to murine FcRn.7 Likewise, the N434Y variant has increased binding to murine FcRn at both pH6.0 and 7.4 resulting in comparable half-life to the IgG-WT in mice.58 Nevertheless, we show in this study that several variants including this mutation N434Y result in half-life extension in hFcRn mice, emphasizing the limitations of using wild-type mice as preclinical models for the analysis of engineered mAbs.

In addition, other factors than FcRn binding have been shown to influence PK of natural mAbs: overall protein structure, glycosylation status, isoelectric point,59 methionine oxidation,59 interaction with the target but also specificity. Indeed, mAbs with the same wild-type human Fc sequences but different Fv domains were shown to bind FcRn with considerable differences.60,61 Conversely, the Fc region has been shown to contribute to affinity and specificity of mAbs.62 For all these reasons, we didn’t focus our study on only one variant but several combinations that are stored and documented in a database, to be able to propose Fc variants that fit best for a given mAb. Preliminary results on two other specificities confirm that several combinations are directly transferable to other mAbs, resulting in increased half-life in hFcRn mice (data not shown).

Finally, for the first time we clearly show that glyco-engineering to improve mAbs functional activity and Fc-mutations to increase half-life can be combined to further optimize mAbs. Beside glyco-engineering, protein engineering has been successfully employed as well to generate antibodies with enhanced binding to activating FcyRs. Alanine scanning,27 in silico approach,63 and random mutagenesis strategy64 allowed identification of distinct IgG1 variants (combinations of 2–5 mutations) with strongly enhanced binding to FcyRIIIA or FcγRIIA resulting in increased ADCC or ADCP in vitro and better anti-tumor activities in mouse models in vivo. We observed that these mutations have little or no effect on FcRn binding (our unpublished data). On the contrary, produced as low fucosylated mAbs, our IgG variants display both advantages: increased ADCC and longer serum persistence. Because we used a random mutagenesis approach, identified mutations could have resulted in a decreased binding to FcγRIIIA. Only one variant, C6A-66 showed poor affinity for this receptor, probably due to an unusual mutation: the deletion of amino acid 294. This variant is nonetheless of great interest as for certain clinical applications no effector function is preferable65,66 and for this variant ADCC is completely abolished while half-life is largely enhanced.

**Materials and Methods**

**Expression and purification of human FcRn**

The expression of soluble human FcRn was performed by GTP Technology (Labège, France) using the baculovirus system as previously described.64 The α chain cDNA encoding the leader peptide and extracellular domains (codons 1–290) was tagged with a TEV sequence and a 6 × polyhistidine tag. The derivative α-chain and the β2-microglobulin chain were cloned into pFastBacDual under the P10 and polyhedrine promoters, respectively. A biotinylated version of FcRn (FcRn-biot) was prepared by chemical coupling with the FluorReporter® Biotin-XX Protein Labeling Kit, F2610 (Molecular Probes) according to the manufacturer’s protocol. A fusion protein containing the β2-microglobulin chain and the α-chain fused to the N-terminal part of the bacteriophage p3 protein and the CVDE protein (FcRn-p3) was also produced. Proteins with more than 90% purity were obtained after IgG-Sepharose and IMAC purification steps.

**Construction of the Fc libraries using Mutagen™**

The human Fc gene encoding amino acid residues 226–447 (EU index as in Kabat), derived from a human IgG1 heavy chain (G1m1.17 allotype), was cloned into the phagemid vector pMG58 (pMG58-Fc226) as a BamHI/EcoRI fragment using standard PCR protocols. Several fully randomized libraries were then generated using the Mutagen™ procedure that uses low fidelity human DNA polymerases (pol. or mutases) to introduce random mutations homogeneously distributed over the whole target sequence. Three distinct mutases (pol. β, pol. η and pol. ι), produced and purified as described previously,64,65 were used in different conditions to create complementary mutational patterns. The human Fc gene was replicated with mutases using the 5′ primer MG-619: 5′-AGTACTGACT CTACCTAAGGA TCCCTGCCCAC CGTGC-3′ and the 3′ primer MG-621: 5′-ACTGCTCGAT GTCCGTACTA TGCGGCCGCG AAATT-3′ (BamHI and EcoRI restriction sites are underlined and italic characters correspond to the non-specific tails). A mixture containing 0.6 μg of the pMG58-Fc226 plasmid as template (wild type Fc for the Mut1 library or Fc variants for the Mut2 library), primers MG-619 and MG-621 (250 nM each) and the appropriate replication buffer (detailed below) was treated for 5 min at 95 °C and immediately cooled down to 4 °C to denature DNA strands. For pol. β, replication buffer was 50 mM TRIS-HCl pH 8.8, 10 mM MgCl₂, 10 mM KCl, 1 mM DTT and 1% (v/v) glycerol. Replication buffer for pol. η (or pol. η and pol. ι) was 25 mM TRIS-HCl pH 7.2, 5 mM MgCl₂, 10 mM KCl, 1 mM DTT, and 2.5% (v/v) glycerol. After
the denaturation step, mutagenic replications were performed by adding 50 μM dATP/dCTP, 100 μM dTTP/dGTP and 1 μg of pol. β or 100 μM dNTPs and 1 μg of pol. η (or pol. η and pol. τ, 1 μg of each mutase). The reaction replication was performed at 37 °C for two hours. The replication products were then desalted and concentrated on Microcon® centrifugal filters (Millipore) before amplification through a selective PCR with tail primers. The primers (MG-619 and MG-621) contain a tail that is non-complementary to the template allowing specific amplification of the DNA fragments synthesized by the mutases. A fraction of the replication products was added to a mixture containing the PCR buffer (20 mM TRIS-HCl pH 8.4, 50 mM KCl), 1.5 mM MgCl₂, 10 pmol of the 5’ and 3’ primers, 200 μM dNTPs and 1.25 U Platinum Taq DNA polymerase (InvitroGen). The PCR cycles were as follows, first cycle: 2 min. at 94 °C, 10 s at 64 °C, 30 s at 7.5 °C, 1 min. at 94 °C, and then 30 selective cycles: 20 s at 94 °C and 30 s at 75 °C. The amplified replication products were purified on 1% (w/v) agarose gels, digested with BamHI and EcoRI restriction enzymes and cloned into the pMG58 vector. The ligation mixtures were used to transform electrocompeptent *E. coli* XL1-Blue cells and subsequently plated on solid 2YT medium supplemented with 100 μg/ml ampicillin and 1% (w/v) glucose. After growth, the number of colonies was determined to estimate the size of the libraries and cells were scrapped in 2YT medium with 15% glycerol, frozen and kept at -80 °C.

Phage display expression of Fc libraries and selection of variants with improved FcRn binding

Fc libraries were expressed on the surface of the bacteriophage M13 using standard procedures. E. coli XL1-Blue bacteria containing the Fc library (pMG58 vector) were grown in 60 ml of 2YT supplemented with 100 μg/ml ampicillin, 15 μg/ml tetracycline and 1% (w/v) glucose at 30 °C. 230 rpm until OD₆₀₀ₙ₉₃ = 0.6 is reached. Cells were then infected with M13 helper phage (M13KO7, New England Biolabs, ratio bacteria: phage = 1:3) at 37 °C for 20 min and phage-Fc production was continued overnight at 26 °C, 230 rpm in 2YT/ampicillin/glucose with 0.5 mM IPTG and 30 μg/ml kanamycin. The following day, phages were precipitated with PEG6000 using standard protocols, resuspended in 1ml phosphate buffer pH6.0 (100 mM sodium phosphate, 50 mM sodium chloride pH6.0, called P6) and titaated by infecting XL1-Blue bacteria. For solid phase replications, 4 × 10¹⁵ phages in P6/5% skimmed milk were incubated into 8 wells of Maxisorp immunoplates previously coated with 0.5 μg neurtavadin and 0.5 μg biotinylated FcRn or 0.5 μg FcRn-p3 and blocked with 5% skimmed milk in P6. After incubation for 2 h at 37 °C, wells were washed 20 times with P6/0.1% Tween-20 and eluted by incubation in 100 μl phosphate buffer pH 7.4 (100 mM sodium phosphate, 50 mM sodium chloride, pH 7.4) per well for 2 h at 37 °C. After titration, eluted phages were used to reinfect 10 ml of exponentially growing XL1-Blue bacteria and subsequently plated on solid 2YT medium/ampicillin/glucose. On the following day, cells were scrapped in 2YT medium with 15% glycerol, frozen and kept at -80 °C until the next round of selection. For liquid phase selection, 4 × 10¹ⁱ phages were first incubated with 250 nM or 100 nM biotinylated FcRn in 1ml P6/5% skimmed milk/0.1% Tween-20 for 1 h at room temperature (RT) under low agitation. Streptavidin-coated magnetic beads (Dynal), previously blocked with 5% skimmed milk in P6 were then added to the phages for 30 min. at RT. Phage-bead complexes were washed 15 times with P6/0.1% Tween-20 using a magnet (magnetic particle concentrator, Dynal). Phages were eluted by incubation in 500 μl phosphate buffer pH 7.4 (100 mM sodium phosphate, 50 mM sodium chloride, pH 7.4) for 2 h at RT. Beads were discarded using the magnet and eluted phages in the supernatants were collected. After titration, eluted phages were used to reinfect 10ml of exponentially growing XL1-Blue bacteria and subsequently plated on solid 2YT medium/ampicillin/glucose. The following day, cells were scrapped in 2YT medium with 15% glycerol, frozen and kept at -80 °C until the next round of selection. During selection processes, for each strategy, from round 3 to round 8, 48 to 96 clones were sequenced as described above. Sequence analysis was performed using MilleGen's proprietary software AnalyseFc internally developed to rapidly analyze the selected Fc variants. Fc variants were named according to the round of selection from which they were isolated. For Mut1 screening: B3A to B6A for solid phase selection on FcRn-p3, S3A to S6A for solid phase selection on biotinylated FcRn and L3A/B to L6A-F for liquid phase selection, for Mut2 screening: C3A to C8A for solid phase selection on FcRn-p3, T3A to T8A for solid phase selection on biotinylated FcRn and M3A/B liquid phase selection. Numbers (1 to 96) refer to the localization of the clone on the PCR plate used for sequencing.

Phage-ELISA assays of the selected variants

The binding characteristics of the variants isolated during selection displayed on the phage were determined using an ELISA test at pH 6.0 with FcRn-p3 coated on wells. Briefly, phage-Fc variants were produced as separate clones on a 96-well plate in 800 μl cultures in 2YT medium/ampicillin/glucose infected with helper phage M13K07 (as described previously). Phages produced overnight at 26 °C were recovered in the supernatants after centrifugation for 30 min at 3000 g. These supernatants were directly diluted (1/2 and 1/4) in P6/5% skimmed milk/0.1% Tween-20 and tested on Maxisorp immunoplates previously coated with 0.25 μg FcRn-p3/well and blocked with 5% skimmed milk in P6. After incubation for 2 h at 37 °C, wells were washed 3 times with P6/0.1% Tween-20 and bound phages were detected with an HRP anti-M13 antibody (GE Healthcare).

Directed mutagenesis

Directed mutagenesis was performed to construct positive controls and remove point mutations using standard PCR protocols. The first positive control: Fc-QL30.31 T250Q/M428L was obtained with two long primers (the 5’ primer 5’-CGGGATCCTG CCCACGGTGCA CCAGCACCGT GAACTCTGGG GGGAGCCTCA GTCTTCTCCT TCCCCCAAAA ACCCAGGGAC CAACTCATGA
TCTCCGGAC-3’ and the 3’ primer 5’-GCAGTTCTTTTACCCGAGAGG CAGGTCTTGCAGTG CAGCGTTCTCCAGTCG
GAGGAGACGC CGGAGCATGA GAAG-3’ and the 3’ primer 5’-CACCTCTACA TCAACCGGAGA ACCTGAGGTCA CACTGCGTGG TG-3’ where overlapping regions are underlined and mutated nucleotides are in italic).

Production and purification of IgG variants

The Fc variants were produced in IgG format with an anti-CD20 specificity (based on the Fv domain of CAT 13.6E1236) using the YB2/0 cell line (ATCC, CRL-1662). For comparative purposes, IgGs based on wild-type Fc (IgG-WT) and Fc-YTE (IgG-YTE) were also produced. To maximize the productivity in the YB2/0 cell line, the full-length light and heavy chain cDNAs including the mutations in the Fc fragment were neo-synthesized with codon optimisation for Rattus norvegicus (GeneArt, Life Technologies). Unwanted features such as cryptic splicing sites or restriction sites were removed. Only one restriction site (ApaI) was present at the junction of variable and constant region. In a first step, wild-type heavy chain was cloned between Nhel and Ascl in the expression vector CHK622-08, optimized for expression in YB2/0, resulting in the intermediate construct HCD20-Opti-GA. The optimized light chain was then cloned between SpeI and XbaI restriction sites resulting in the final construct HKCD20-Opti-GA for expression of the IgG-WT. Fc variants were prepared by replacing the wild-type IgG1 Fc fragment present in HKCD20-Opti-GA by the different mutant versions which were cloned between ApaI and Ascl restriction sites. These cloning steps were done by classical digestion/ligation procedures, followed by bacterial transformation. Expression constructs were screened by enzymatic digestion and PCR and validated by sequencing. Each linearized expression construct was introduced by electroporation into 5.10^6 YB2/0 cells. Cells were then diluted to 25,000 cells/ml in RPMI 1640 medium + 5% v/v dialysed FCS (Invitrogen) and dispensed in 1 ml/well within 24-well plates. After 3 d, selection pressure was applied by adding gentamicin (Invitrogen) and methotrexate (Sigma) to obtain final concentrations of 0.5g/l and 25 mM respectively, in 2ml/well. After 11 d, resistant cells were pooled for each construct (encoding for the selected IgG variants, IgG-YTE and IgG-WT) and progressively diluted with DMEM medium + 5% v/v Ultra-low IgG FCS (Invitrogen) until two 2 L-roller bottles, each containing 0.9 L of cell suspension, can be incubated at 2rpm. Cells were allowed to grow and die (4 to 5 d) before supernatant collection, clarification by low-speed centrifugation and volume reduction by ultra-filtration on Pallion XL Filter (Millipore). The concentrated culture supernatants were injected into a HiTrap protein A FF column (GE Healthcare). Bound antibodies were eluted with 0.1 M sodium citrate, pH 3.0 and fractions were neutralized using 100 μl of 1M TRIS-HCl pH 7.5 per ml of elution buffer. Fractions containing the antibodies were pooled and dialyzed against PBS pH 6.0, and the samples were sterile-filtered (0.22 μm) and stored at 4 °C. The purified IgGs were analyzed using SDS-PAGE under non-reducing and reducing conditions. Coomassie Blue-stained gels indicated that the IgGs, whatever the mutations, had greater than 95% purity and displayed the characteristic heavy and light chain bands. Low aggregate levels and absence of contaminants were verified by analytical gel filtration in PBS on superdex 200 10/300 GL with an AKTA prime system (GE Healthcare). Purified IgGs were considered suitable for biological tests when aggregate rates were below 5%. None of the mutations tested induced IgG aggregation, aggregation rates of the purified IgGs were between 0.1 and 4.5%. Final IgG concentrations were between 3.5 and 8.2mg/ml. LAL endotoxin test (Limulus Amebocyte Lysate) gel clot method was further used to test purified IgGs for the presence of endotoxins. Endotoxin levels of the purified IgGs were below 7 UI/mg.

FcRn binding of the IgG variants

Three distinct tests were set up to characterize the interaction of the IgG variants with FcRn: (1) an ELISA test, (2) a competitive binding assays on FcRn expressing Jurkat cells, and (3) SPR. (1) The binding properties of the IgG variants, in comparison with the IgG-WT, were determined using an ELISA test at pH 6.0 with FcRn-p3 coated on wells. Purified IgG variants serially diluted in P6/5% skimmed milk/0.1% Tween-20 were tested on Maxisorp immunoplates previously coated with 0.1 μg FcRn-p3/well and blocked with 5% skimmed milk in P6. After incubation for 2 h at 37 °C, wells were washed 3 times with P6/0.1% Tween-20 and bound IgG variants were detected with an HRP F(ab’)2 goat anti-human F(ab’)2 (Interchim). For each IgG variant, the percentage of bound FcRn was plotted vs. the log of the concentration of IgG variant. For each resulting binding curve, the IgG concentration corresponding to 50% saturation of the curve (EC50) was determined and compared with the EC50 of IgG-WT. (2) Competitive immunofluorescence assays were performed to evaluate the ability of IgG-WT and IgG variants to interact with FcRn by a method adapted from that described elsewhere. Briefly, IgG-WT and IgG variants were diluted in PBS pH6.0 at a final concentration ranging from 0.06 to 2 mg/ml and incubated with 1.5 x 10^5 Jurkat cells expressing on the cell membrane a truncated form of FcRn in reducing conditions. Coomassie Blue-stained gels indicated that the IgGs, whatever the mutations, had greater than 95% purity and displayed the characteristic heavy and light chain bands. Low aggregate levels and absence of contaminants were verified by analytical gel filtration in PBS on superdex 200 10/300 GL with an AKTA prime system (GE Healthcare). Purified IgGs were considered suitable for biological tests when aggregate rates were below 5%. None of the mutations tested induced IgG aggregation, aggregation rates of the purified IgGs were between 0.1 and 4.5%. Final IgG concentrations were between 3.5 and 8.2mg/ml. LAL endotoxin test (Limulus Amebocyte Lysate) gel clot method was further used to test purified IgGs for the presence of endotoxins. Endotoxin levels of the purified IgGs were below 7 UI/mg.
a BIAcore X100 instrument using a CM5 sensor chip (BIAcore, GE Healthcare). The methodology was similar to that previously described for analyzing Fe-FcRn interactions. Recombinant soluble FcRn was coupled to flow cell 2 of the sensor chip using amine-coupling chemistry. The flow cells were activated for 3 min with a 1:1 mixture of 0.1M N-hydroxysuccinimide and 0.1M 3-(N,N-dimethylamino)propyl-N-ethylcarbodiimide at a flow rate of 30 μl/min. Recombinant human FcRn (5 μg/ml in 10 mM sodium acetate, pH 5.0) was injected over flow cell 2 for 8 min at 10μl/min, which resulted in a surface density of 1200 to 1300 response units (RU). Surfaces were blocked with a 3 min injection of 1 M ethanolamine-HCl, pH 8.5. Flow cell 1 was used as a control surface without FcRn and was prepared similarly to sample flow cell. The data from this blank flow cell were subtracted from the sample data. IgGs were diluted in PBS/Tween-20 (50 mM sodium phosphate, pH 6.0, 150 mM NaCl, 0.02% NaN₃, 0.01% Tween-20) which is used as running buffer in equilibrium binding experiments. All measurements were performed at 25 °C with IgG concentrations typically ranging from 1 to 200nM at a flow rate of 10 μl/min. Data were collected for 10 min. and a 1 min pulse of PBS, pH 8 containing 0.05% Tween-20 was used to regenerate the surfaces. Sensorgrams were generated and analyzed by using the steady-state affinity model included in BIAevaluation software version 3.1. The equilibrium RU observed for each injection was plotted against the IgG concentration. The equilibrium Kₘ values were determined using the steady-state affinity model included in the BIAevaluation software.

**hFcRnIII A binding of the IgG variants**

The human recombinant FcyRIIIA (F158 allotype) made by PX'Therapeutics was biotinylated with the EZ-link NHS-PEO kit (Pierce), diluted to 1 μg/ml in assay buffer (25 mM Tris, 150 mM NaCl, pH 7.35, 0.05% Tween-20, 0.1% BSA) and coated onto react-bind™ streptavidin ELISA plates (Pierce) for 2 h at RT. During this incubation time, IgG-F(ab')₂ complexes were prepared in assay buffer by mixing 5 μg/ml of IgG and 2 μg/ml F(ab')₂, anti-human F(ab')₂ labeled with horseradish peroxidase (Jackson ImmunoResearch) for 2 h at RT. Serial dilutions of complexes were added to plates and incubated for 2 h at RT under gentle shaking. After washing plates with assay buffer, bound complexes to hFcRnRIIIA were detected with TMB (Pierce). Absorbance at 450 nm was read using a plate reader (Tecan). For each IgG variant, the percentage of bound hFcRnRIIIA (which is obtained from OD450nm) was plotted against the concentration of IgG variants.

**ADCC activity**

Natural killer cells (NK cells) were purified from the peripheral blood of healthy volunteer donors by the negative depletion technique developed by Miltenyi. The ADCC test comprises incubating the NK cells with CD20-expressing Raji target cells, in the presence of different concentrations of anti-CD20 antibodies. After 16 h of incubation, the cytotoxicity induced by the anti-CD20 antibodies was measured by quantifying in cell supernatants the level of the intracellular enzyme lactate dehydrogenase (LDH) released from lysed target cells. The specific lysis results are expressed as the percentage of lysis as a function of antibody concentration. EC₅₀ values (antibody concentration inducing 50% of maximum lysis) were calculated using PRISM software.

**PK study in human FcRn mice (hFcRn mice)**

Animals were housed in the animal center of the Faculty of Pharmacy, University of Paris-Sud 11 and all studies were approved by the responsible Animal Care and Use Committee (authorization number 2012_121, CEEA 26). hFcRn mice (mFcRn⁻/⁻ hFcRnTg 276 heterozygote on a B6 background) for PK studies were obtained by breeding of homozygous Tg276 human FcRn mice with FcRn⁺ mice obtained from The Jackson Laboratory. These mice are deficient in the mouse FcRn α-chain and carry a human FcRn α-chain gene instead. For PK studies, each animal (4 animals per group) received a single dose of 5 mg/kg of antibody intravenously via the retro-orbital sinus. Blood samples were collected from the retro-orbital sinus at the following time points: 1 h, 4 h, 6 h, 24 h, 48 h, 96 h, 168 h, 216 h, 264 h, 336 h, 384 h, 432 h, and 504 h, and transferred into tubes. Three to five mice per group underwent retro-orbital bleeding for each time point, with 100 μl of blood collected per mouse (5 punctures distributed over 3 wk for each mouse). The blood samples were left at RT for at least 1 h, processed to serum and stored at -80 °C until analysis. A qualified anti-human IgG immunoassay was used to determine all mAb levels in the serum samples and was performed by LFB biotechnologies or Bertin Pharma. Briefly, an affiniPure Donkey Anti-Human IgG (H’L) polyclonal antibody was used for capture, and an affiniPure Donkey Anti-Human IgG (H’L) peroxidase polyclonal antibody was used for detection (both from Jackson ImmunoResearch Laboratories). The mAb concentrations were extrapolated from standard curves. Samples were diluted at least 40-fold. The assay range was established with spiked quality control samples. The lower limit of quantification was 60 ng/ml. Sample results were corrected for dilution; values within the standard curve range were averaged. The serum concentration was used for non-compartmental PK analysis using WinNonlin (Enterprise version 5.01). The mAb elimination phase terminal half-life (t₁/₂) was determined using data points from the terminal phase. These data points generally fitted well to a monoexponential decay function.

**Disclosure of Potential Conflicts of Interest**

CM, SJ, AJ, NF, CDR, RU, CKB, PM, and AF are employees of LFB Biotechnologies and NS, OZ, FC and KB are employees of MilleGen; both companies financially supported the study.

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