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FcγRIIib on Liver Sinusoidal Endothelium Clears Small Immune Complexes

Latha P. Ganesan,* Jonghan Kim,† Yun Wu,‡ Sudhasri Mohanty,* Gary S. Phillips,§ Daniel J. Birmingham,* John M. Robinson,¶ and Clark L. Anderson*†

It has long been known that the ITIM-bearing IgG Fc receptor (FcγRIIib, RIIb) is expressed on liver sinusoidal endothelial cells (LSEC) and that the liver is the major site of small immune complex (SIC) clearance. Thus, we proposed that RIIb of LSEC eliminates blood-borne SIC, thereby controlling immune complex-mediated autoimmune disease. Testing this hypothesis, we found most RIIb of the mouse, fully three-quarters, to be expressed in liver. Moreover, most (90%) liver RIIb was expressed in LSEC, the remainder in Kupffer cells. An absent FcγR in LSEC implied that RIIb is the sole FcγR expressed. Testing the capacity of liver RIIb to clear blood-borne SIC, we infused mice intravenously with radio-iodinated SIC made of OVA and rabbit IgG anti-OVA. Tracking decay of SIC from the blood, we found the RIIb knockout strain to be severely deficient in eliminating SIC compared with the wild-type strain, terminal half-lives being 6 and 1.5 h, respectively. RIIb on LSEC, a major scavenger, keeps SIC blood concentrations low and minimizes pathologic deposition of inflammatory immune complex. The Journal of Immunology, 2012, 189: 4981–4988.

It is an axiomatic function of the immune system that complexes of Ag and Ab are eliminated efficiently from the bloodstream, ridding the body of harmful elements of a variety of sorts, both invasive and endogenous. A number of features of immune complexes govern their elimination, the most obvious being size. Large immune complexes composed of, for example, bacteria-size microorganisms and blood cells coated with Ab are removed by cells of the mononuclear phagocyte system by the process of receptor-mediated phagocytosis. These complexes are sometimes referred to as “insoluble” because they readily sediment at low gravity forces.

Likewise, small immune complexes (SIC), the size of serum proteins complexes with corresponding Ab, are in most situations eliminated efficiently from the blood, specifically by the cells of the classical reticuloendothelial system, chiefly by the sinusoidal cells of the liver (1–4). These SIC are often termed “soluble” because they fail to sediment at low gravity forces; experimentally, they are prepared at “antigen excess,” at concentrations of Ag exceeding that required for the optimal preparation of insoluble immune complexes (5). In certain circumstances, however, these SIC fail to be eliminated and go on to produce disease. The pathophysiology of such immune complex-mediated diseases has been the focus of considerable experimental interest over the past century. Intensive study of the mechanisms of these diseases, beginning with experimental models of serum sickness in the 1950s most prominently by Dixon and colleagues (6), Benacerraf and colleagues (7), and others, has led to the current concept that SIC must accrete to a certain requisite size to deposit in critical organs, such as the kidney, where they produce inflammation and disease (8–11). Thus, analyzing the mechanism by which immune complexes, while still small, are eliminated by the liver becomes an imperative theoretical and practical objective.

A crucial clue to the mechanism of SIC elimination emerged 30 y ago when it was observed that mab 2.4G2, specific for the binding sites of all mouse FcγR, blocked liver uptake of SIC infused intravenously into mice (3). This finding implicated liver sinusoidal FcγR in the elimination of SIC. Many concluded that the Kupffer cells (KC) were responsible for such liver elimination. Rather, as we have recently found, liver sinusoidal endothelial cells (LSEC) expressing FcγRIIb (RIIb) are responsible for virtually all mab 2.4G2 binding in mouse liver (12). Several converging lines of recent evidence (4, 12–18) now indicate that the predominant FcγR on LSEC is RIIb, the low-affinity ITIM-bearing inhibitory receptor studied most carefully on B cells, macrophages, and dendritic cells (reviewed in Refs. 19, 20).

We are led by the evidence presented above to propose that RIIb expressed on LSEC are responsible for the ongoing removal of SIC from the blood. This hypothesis has two predictions; first, that liver expression of RIIb is remarkably abundant; second, that a mouse strain lacking RIIb will fail to remove SIC from blood. We tested these two predictions, the first by immunofluorescence microscopy and immunoblotting, measuring the expression of RIIb in liver compared with the entire animal; and the second by comparing the rates of elimination of radio-iodinated SIC from the blood in wild-type (WT) and RIIb knockout (KO) strains of mice.

Neither test rejected the hypothesis. Rather, a very high fraction, fully three-quarters, of total body RIIb was found in the liver, most
on LSEC. Further, the removal rate of SIC from blood of RIIb KO mice compared with WT mice was severely curtailed. These results confirm our hypothesis and suggest new ways of conceptualizing and treating immune complex-mediated diseases.

Materials and Methods

Ethics statement and animals

Male mice of age 12–15 wk were obtained from Taconic Laboratory. They were of strains BALB/c WT and RIIb KO on a BALB/c background [C.129S4 (B6)-FcyRIIbJtm1Tg6KcAnNTac N12C.129S4 and model number 580-M]. The RIIb KO mice were generated and described previously (21). All protocols were approved by The Ohio State University Institutional Animal Care and Use Committee. Bleedings were performed under isoflurane anesthesia, and all efforts were made to minimize suffering.

Cells

RAW 264.7, A20, and COS-7 cells were obtained from American Type Culture Collection. COS-7 cells were maintained in DMEM supplemented with 10% FBS. RAW 264.7 and A20 cells were maintained in RPMI 1640 medium supplemented with 5% FBS. Bone marrow macrophages (BMM) were obtained and differentiated with M-CSF as previously described (22).

Quantitative immunoblot

The expression of RIIb and Syk in various organs was quantified by immunoblot as follows: organs (kidney, spleen, ileum, liver, heart, and lung) from three sacrificed mice were harvested and weighed. Small pieces of 100 μg from each organ were homogenized with a glass homogenizer in a lysis buffer composed of 25 mM HEPES, 20 mM Na4H2O7·10H2O, 100 mM NaF, 4 mM EDTA, 2 mM Na3VO4, 1% Triton X-100, 0.34 mg/ml PMSF, 0.01 mg/ml aprotinin, and 0.01 mg/ml leupeptin. Lysates were loaded onto nitrocellulose membranes by 0.45 μm. The membranes were blocked after washing with PBS. Bone marrow cells were flushed from femurs, centrifuged at 376 × x for 5 min, and lysed. Blood (∼80 μl) was obtained from mice via the retroorbital plexus using heparinized capillaries, and the erythrocytes were lysed by dilution in 10 vol of 20% PBS at room temperature for 10 min. Cells remaining were spun down at 376 × x for 5 min and lysed. All lysates were incubated on ice for 30 min and centrifuged at 18,407 × g for 15 min. The protein concentrations of spun lysates were estimated using the bicinchoninic acid protein assay.

Organ lysates (35 μg) along with lysates of control standards (0.3, 6, 9, 12 μg of lysates from A20 cell line for RIIb blots and 0.5, 10, 15, 20 μg of lysates from RAW 264.7 cells for Syk blots) were separated on 8–16% gradient SDS-polyacrylamide gels, and the proteins were transferred to nitrocellulose membranes (0.45 μm). The membranes were blocked with 5% milk for 30 min and then incubated overnight with primary rabbit anti-mouse RIIb (kind gift from John Cambier) and rabbit anti-Syk (Santa Cruz Biotechnology), and rabbit anti–FcγR-IIb (Millipore) were all used at dilutions of 1:25. The membrane was used at a concentration of 20 μg/ml. After three washes with PBS, the sections were incubated with secondary Abs for 1 h at room temperature. Nuclei were stained with DAPI for 10 min, and the sections were mounted under coverslips in Prolong gold (Invitrogen). Controls incubations included isotype controls along with their respective secondary Abs and also secondary Abs alone. The images were acquired in the Olympus FV1000 Laser Scanning Confocal Microscope equipped with a spectral detection system for a finer separation of fluorochromes (FV 1000 spectra) using a ×60 oil immersion lens at room temperature. Image analyses were done using Fluoview software (Olympus version 2.1.39). In Fig. 4, to overcome the inherent limits of resolution in the z-dimension in our confocal microscopes, we used ultrathin cryosections of liver tissue as the substrate for high-resolution immunofluorescence microscopy as we have described (27–29). Preparation of tissue and ultrathin cryosections were described in detail earlier (30).

Quantitative microscopy

The relative liver expression of RIIb between LSEC and KC, shown in Fig. 3 (bottom row), was quantified from immunofluorescence images dual-stained for RIIb (green) using goat anti-RIIb Ab and anti-F4/80 (red) in the following manner. The total intensity of RIIb (green) in the images was calculated using ImageJ software. From this total was subtracted the intensity of RIIb measured in all cropped KC, identified with F4/80. Thus, total intensity of RIIb minus KC intensity of RIIb equaled the intensity of RIIb in KC. Optical sections with a total area of 6.3 mm2 were analyzed from each of three mice. The fluorescence intensities from mannose receptor (MR) (green) and RIIb (red) channels in the ultrathin sections represented in Fig. 4 were obtained using a ×510 META software. Randomly selected sinusoids (n = 20) from merged color images were cross-sectioned eight times at 45˚ angles, and the intensities of both color channels of eight sections over the LSEC cross-sectional distance from luminal to abluminal surface were recorded. Data were analyzed using a paired t test.

Immunofluorescence from anti–FcγR-IIb chain (green) was colocalized with signals from mab F4/80 and mab 2.4G2 (Fig. 6) in 60 immunofluorescence optical sections with an area of 12.6 mm2 from a total of three mice. Using Olympus FV10-ASW analysis software, the Mander’s overlap coefficient R was obtained and expressed as a percentage of colocalization (31).

Immunofluorescence signal from mab 2.4G2 colocalized with signal from rabbit anti-RIIb shown in the top row of Fig. 2 was quantified as described for Fig. 6, with the following modification. Individual LSEC (n = 60) were cropped from images obtained from two WT livers, and Mander’s overlap coefficient was expressed as percent colocalization. By cropping the LSEC, we avoided pixels from nuclei nonspecifically stained with the rabbit RIIb Ab. Signal from mab 2.4G2 colocalization with immune fluorescence from goat anti-RIIb in individual LSEC (n = 194) was quantified manually by scoring the number of LSEC that were positive for 2.4G2 (green) and goat anti-RIIb (red) (Fig. 2, bottom row).

RT-PCR

Total RNA extracted from TRZol lysates of organs and cell lines were transcribed to cDNA using Thermoscript RNaScript H−reverse transcriptase and amplified using primers specific for mouse RIIb. The RNA isolation protocol, primer sequences, and PCR conditions were described earlier (32). The RT-PCR products were resolved by electrophoresis on 1.5% agarose gels, stained with ethidium bromide, and imaged using Quantity One software.

Immuno fluorescence

Small pieces of liver (∼5 mm) were fixed in 4% paraformaldehyde in PBS for 2 h at room temperature and, after washing with PBS, were infused in 20% sucrose–PBS overnight at 4˚C. The tissues were then embedded in a freezing medium and stored at −80˚C. Cryostat sections, 5 μm thickness, were blocked in 5% milk–PBS prior to incubation overnight at 4˚C with primary Abs. The primary Abs rabbit anti-mouse RIIb, goat anti-mouse RIIb (kind gift from Mark Coggleshall), rabbit anti–mannose receptor (Santa Cruz Biotechnology), and rabbit anti–FcγR-IIb (Millipore) were all used at dilutions of 1:25. The rabbit 2.4G2 was used at a concentration of 20 μg/ml. After three washes with PBS, the sections were incubated with secondary Abs for 1 h at room temperature. Nuclei were stained with DAPI for 10 min, and the sections were mounted under coverslips in Prolong gold (Invitrogen). Controls incubations included isotype controls along with their respective secondary Abs and also secondary Abs alone. The images were acquired in the Olympus FV1000 Laser Scanning Confocal Microscope equipped with a spectral detection system for a finer separation of fluorochromes (FV 1000 spectra) using a ×60 oil immersion lens at room temperature. Image analyses were done using Fluoview software (Olympus version 2.1.39). In Fig. 4, to overcome the inherent limits of resolution in the z-dimension in our confocal microscopes, we used ultrathin cryosections of liver tissue as the substrate for high-resolution immunofluorescence microscopy as we have described (27–29). Preparation of tissue and ultrathin cryosections were described in detail earlier (30).

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ENDOTHELIAL FcγRIIb CLEAR IMMUNE COMPLEXES
A single lot of affinity-purified rabbit IgG anti-OVA Ab (Immunology Consultants) was “calibrated” for its interaction with OVA (Sigma) by a quantitative precipitin reaction to derive a point of equivalence (POE) (2, 5, 33). Specifically, microfuge tubes containing serially diluted OVA at concentrations ranging from 6.5 to 15 µg/ml were each incubated with 200 mCi Na¹²⁵I-labeled OVA (Pierce) in a total volume of 500 µl of PBS at pH 7.4 for 1 h at room temperature and 18 h at 4°C. The mixtures were centrifuged at 2000 × g for 10 min; the pellets were dissolved in saline HCl buffer (0.05 M glycine and 0.15 M NaCl pH 2.3); and the protein concentrations of pellets and supernatants were determined by UV absorbance at 280 nm.

The calibration curve of the precipitin reaction yielded a distinct POE at Ag and Ab concentrations of 26 and 200 µg/ml, respectively. To prepare SIC for determination of particle size by dynamic light scattering (DLS) and for infusion into mice, we increased the protein concentrations 5-fold greater than the concentration used in the precipitin curve. SIC were prepared by mixing rabbit anti-OVA Ab at 1000 µg/ml (~5 × 200) with OVA (Ag) at 1950 µg/ml (26 × 5 × 15), a 15-fold excess over the POE. Ag concentration determined in the calibration curve, attempting to duplicate the conditions used by others (2, 3, 7). The mixture was incubated for 1 h at room temperature and 18 h at 4°C and was centrifuged at 2000 × g for 1 min. SIC size was measured by DLS. To prepare trace-labeled SIC for infusion into mice, rabbit anti-OVA Ab (iodinated as described later) was added to unlabeled Ab at a 1:500 mass ratio prior to mixing Ag with Ab (exactly as described earlier).

Radioiodination of Ab
Affinity-purified rabbit IgG anti-OVA Ab was dialyzed against PBS pH 7.4 containing 1.06 mM KH₂PO₄, 155 mM NaCl, and 2.97 mM Na₂HPO₄·7H₂O and radio-iodinated by a modified chloroglycouril method using chloroglycouril-coated tubes (Pierce). Briefly, at room temperature 1 µCi of Na¹²⁵I (Pierce) was activated in a chloroglycouril-coated tube for 6 min, then centrifuged at 2000 × g for 10 min; the pellets were dissolved in glycine HCl buffer (0.05 M glycine and 0.15 M NaCl pH 2.3); and the protein concentrations of pellets and supernatants were determined by UV absorbance at 280 nm.

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Analysis of SIC using DLS
The particle size of SIC was estimated by DLS (BI-200SM Research Goniometer System; Brookhaven Instrument, Holtsville, NY) under the following conditions: detection angle 90°, laser wavelength 633 nm, temperature 20°C, measurement duration 2 min. The particles were diluted in 0.22-µm-filtered PBS. Fe₃O₄ particles of size range 200–250 nm (BD Biosciences), 10, and 5 nm (NN Laboratories) were analyzed to check the precision of the instrument and were found to have mean diameters of 209, 12, and 20 nm, respectively, suggesting imprecision at low particle size. The standard latex particles of size 40, 92, and 100 nm were found to have mean diameters of 42, 93, and 98 nm, respectively. The mean diameters by volume of Ag and Ab at a concentration of 1 mg/ml were 6 ± 1 and 5 ± 1 nm, mean ± variance of two observations, respectively. The mean diameter by volume of particles at the POE was 1228 ± 774 nm. The mean diameter by volume of SIC particles prepared at 3-fold excess Ag was measured to be 220 ± 14 nm. The characterization of SIC prepared at 15-fold excess Ag is presented in Fig. 1B and the text. All samples showed a monodispersed distribution with insignificant outliers and aggregates.

Clearance kinetics of SIC
To characterize in vivo SIC clearance kinetics, we infused by tail vein freshly prepared radio-iodinated SIC containing 1.4 × 10⁶ and 1.9 × 10⁶ cpm in 58 µl. Two identical experiments were performed 3 mo apart, in each infusion three WT and three RIIb KO mice matched for sex and age. The mice were bled of ~15 µl blood via the retro-portal plexus at post-infusion times of 1, 5, 10, 20, 30, and 60 min. To adjust for different body weights of individual mice, the blood concentrations of radioactive SIC (cpm/10 µl blood) were normalized to an average value of dose/body weight among animals (i.e., a dose of 1.6 × 10⁶ cpm/25.43 g body weight). Because a semilog plot of blood concentrations of iodinated SIC from both strains indicated biphasic decay, we used a biexponential decay model to fit the SIC concentration–time profile. Half-lives were calculated by 0.693/∼slope, where the slope was obtained from the biexponential decay analysis. Independent two-sample t-test was performed to compare the means of two genotypes. Differences between WT and KO strains were considered significant at p < 0.05.

Quantification of SIC in various organs
Mice (n = 3) were infused intravenously with freshly prepared radio-iodinated SIC containing 1.1 × 10⁶ cpm in 58 µl and were sacrificed at 25 min. The mice were bled of ~20 µl via the retro-portal plexus; organs (liver, kidney, lung, spleen, and heart) were removed and weighed. SIC in the weighed portions of each organ were measured and quantified after factoring total organ weight.

Results
Most RIIb of the mouse is in the liver
We recently noticed that the expression of RIIb on LSEC was astonishingly high, far in excess of what we had perceived to be expressed on other RIIb reservoirs of the body such as spleen, lymph nodes, B cells, and macrophages (12). More precisely, assessing in our mind’s eye the brightness and extent of RIIb fluorescence in a microscopic field of view of liver sections, and multiplying this by the three dimensions of the liver, the body’s largest internal organ, the total quantity of RIIb would be far greater than that of any of the other RIIb sites of the body (data not shown). It has long been known that a like assessment of the human liver with a specific anti-RIIb Ab gives the identical impression by immunolabeling (14). Quantifying our visual impression, we measured by immunoblotting the expression of RIIb in lyses of the liver and spleen and most of the other major organs and tissues of the body (Fig. 1). Consistent with our visual impression, we found that of the total body pool of RIIb, fully 72 ± 5% (n = 3) was expressed in the liver, and the remaining 28% was spread among the other organs and tissues of the body, each being less than 10%. By mobility of the anti-RIIb identified bands, the liver appeared to express mostly the b2 protein isoform but also some b1, whereas, as expected, most of the RIIb in spleen was b1 (Fig. 1A). Our immunoblots of organ lyses affirm that this RIIb KO strain shows no evidence for the RIIb bands (data not shown). The band with b1 mobility in the kidney lane, as well as unidentified bands at 75 and 45 kDa in many lanes, were artifacts based on their presence in tissues from RIIb KO mice (data not shown). Further studies of kidney tissue sections examined by immunofluorescence (IF) microscopy with three anti-RIIb Abs showed no evidence for the expression of RIIb in the kidneys (data not shown).

Because our method of quantifying tissue expression of proteins from band densities in immunoblots entailed a large multiplicative factor for liver due to its relative size, we also measured the expression of the tyrosine kinase Syk, reasoning that a molecule expressed largely in the spleen would appear highly expressed in liver if our method were artificually amplifying the extent of liver expression (34). Confirming the validity of our method, 63 ± 6% of total body Syk was expressed in the spleen but less than 10% in liver and all other organs except the ileum (13%) where B cells and macrophages are prevalent (Fig. 1B).

LSEC expression of RIIb confirmed with three anti-RIIb Abs
Within the liver, RIIb is predominantly expressed in LSEC (12, 14, 17, 35). We confirmed this conclusion by IF microscopy using three anti-RIIb Abs; that is, mab 2.4G2 and two polyclonal Abs from goat and rabbit, the last directed toward the cytoplasmic tail sequence of RIIb. We found that all three Abs gave similar and mostly overlapping signals in LSEC (Fig. 2); these signals were specific for RIIb in that liver sections from RIIb KO mice showed...
Within the liver, most RIIb is in LSEC

We quantified by IF microscopy of liver sections the relative expression of RIIb between LSEC and KC by colocalizing the binding of two anti-RIIb Abs with the KC marker F4/80 (Fig. 3). We found that of the total liver RIIb pixel intensity, 90% was expressed in LSEC and 10% in KC.

**RIIb expression in LSEC predominates toward the apical membrane**

Having earlier found that RIIb in the endothelium of the human placenta is expressed in the interior of the cell and not at the plasma membrane (28), we used the same strategy to localize RIIb in mouse LSEC, examining ultrathin (<100 nm) sections of liver by three-color IF microscopy, comparing anti-MR binding, which is relatively specific for LSEC (12), with anti-RIIb Ab binding (Fig. 4A). Quantifying the pixels of the two colors in merged images (Fig. 4B), we concluded that, unlike its localization in human placental endothelium, RIIb does not predominate in the interior of the cell; rather it appeared to be more highly expressed toward the apical portion of the LSEC, whereas MR was more diffusely spread throughout the cell.

**The RIIb mRNA isoform expressed in liver is b2**

Having found that LSEC-expressed RIIb constitutes nearly all RIIb of liver (Fig. 3), we realized that we could easily determine the mRNA isoform of RIIb in LSEC by RT-PCR analysis of whole-liver mRNA, comparing band sizes with known sizes from RIIb-expressing cells characterized earlier by others. Such an experiment indicated that the predominant isotype in liver was b2 with only a trace band of b1 being seen in the agarose gels (Fig. 5). We have analyzed the published cDNA sequence of rat RIIb from LSEC that was not isotyped in the original publication and find that the isotype was b2 and not b1 (17). Thus, human, rat, and mouse RIIb, which are highly expressed in LSEC, are of the b2 isotype (14, 16, 17).

**Fcγ-chain is expressed only in KC and not in LSEC**

We were unable to determine with confidence whether other members of the FcγR family were expressed in LSEC because of a dearth of reliable Abs to RI, RIII, and RIV that would be useful in IF microscopy. However, as all three of these agonist receptors require for function the association of the Fcγ-chain (36–38), we realized that by assessing FcγR expression in LSEC, we could...
indirectly assess the expression of the agonist members of the FcγR family in mouse. By three-color IF microscopy, we found FcRγ to be expressed only in the KC of the liver and not in the LSEC (Fig. 6A). Quantifying the colored signals in these images, we affirmed that FcRγ was a distinctive feature of KC and not LSEC (Fig. 6B). Thus, it seems likely that RI, RIII, and RIV are not expressed in LSEC.

Quantitative precipitin curve and SIC characterization

We prepared SIC using a quantitative precipitin curve described 76 y ago by Heidelberger and Kendall (5). Calibrating a single lot of affinity-purified rabbit IgG anti-OVA Ab, we determined the concentrations of Ab and OVA at the POE where the immune complexes precipitated maximally (Fig. 7A). We then prepared SIC at a concentration of Ag 15-fold greater than the POE concentration, conditions found by others to yield SIC (2, 3, 5). Measured by DLS, these SIC had a mean diameter by volume of ∼39 nm with a range from 20 to 150 nm (Fig. 7B).

SIC are not cleared efficiently in the RIIb KO mouse strain

Testing a prediction of our central hypothesis, that the RIIb of LSEC serves the purpose of removing SIC from circulation, we measured the elimination rate of SIC from the bloodstream of both WT and RIIb KO strains. Mice were infused by tail vein with trace radio-iodinated SIC prepared at 15 times Ag excess, as described earlier; the decay of gamma-radiation from peripheral

FIGURE 3. Within the liver, most RIIb is in LSEC. Confocal IF images in the left column show the binding pattern of rabbit anti-mouse RIIb in the top panel and goat anti-mouse RIIb in the bottom panel (both green). In the middle column, both panels show KC whereabouts with anti-F4/80 (red). The merged color images in the right column illustrate the relative abundance of RIIb expression in LSEC versus KC and show also DIC and DAPI staining of nuclei. Scale bars, 10 μm.

FIGURE 4. LSEC RIIb is mostly in the membrane. (A) Two-color IF images of an ultrathin (100 μm) cryosection through mouse liver locating the binding of anti-MR (green, upper left) and anti-RIIb (red, upper right). The bottom panels show merged images of the upper two panels, and the right lower panel adds DIC and DAPI staining. Scale bar, 10 μm. (B) Quantification of relative topological distribution of RIIb and MR in LSEC. The bar graph plots the fluorescence intensities (mean ± SD) of the two colors in LSEC cross-sections drawn from luminal to abluminal surfaces. n = 100 sinusoids. *p < 0.05.

FIGURE 5. The RIIb isoform expressed in liver is b2. Ethidium bromide-stained agarose gel of electrophoresed RT-PCR products using primers specific for RIIb showing the mRNA expression of the two RIIb isoforms in organs and cells including spleen, liver, A20, RAW 264.7, COS-7, M-CSF–differentiated BMM, and b1 cDNA. The two bands at 313 and 172 bp correspond to the two RIIb transcripts b1 and b2, respectively. The data are representative of samples from two sets of WT organs and cell line preparations.
blood was followed over the course of 1 h; and the data in cpm were plotted in semilogarithmic fashion (Fig. 8). Indeed, our prediction was not falsified. Rather, we found that the rate of SIC clearance from the RIIb KO strain was dramatically retarded compared with the WT strain. Specifically, both curves described biexponential decay with early and late phases. Decay was most obvious during the late phase, from 10 to 60 min, where half-lives were 87 ± 20 min and 367 ± 222 min for WT and KO mice, respectively (mean ± SD, \( n = 5 \) to 6, \( p = 0.0001 \)), indicating a 3-fold lengthier survival of SIC in the absence of RIIb. In contrast, decay rates during the early 1- to 5-min period were not statistically different between the two strains, half-lives for WT and RIIb KO strains being 12 ± 6 min and 19 ± 10 min, respectively (\( n = 5 \) to 6, \( p = 0.2 \)) (Fig. 8). Note also that the cpm at 1 min were >20% lower in WT compared with the KO strain (see Discussion).

Quantification of SIC in various organs
Documenting that infused SIC homes mostly to liver, we autopsied three mice 25 min after infusion of radio-iodinated SIC. Of the total administered dose, we recovered 72 ± 2% in blood, 27 ± 2% in liver, 0 ± 1% in lung, 1 ± 0% in spleen, and 0 ± 1% in kidney (mean ± SD). Thus, nearly all of the cleared SIC (96%) was found in the liver.

Discussion
Our results allow two major conclusions. First, RIIb is highly expressed on the endothelial cells of the liver sinusoids. This impression is easily reached by simple inspection of a section of liver stained with anti-RIIb Ab. But, wanting to quantify our visual impression, we have measured by immunoblotting the amount of RIIb expressed in the major organs of the body and have concluded that

![Image 6](https://example.com/image6)

**FIGURE 6.** Fc\(\gamma\)RIIb-chain is expressed only in KC and not in LSEC. (A) The confocal IF image in the left column illustrates the expression pattern of \(\gamma\)-chain (green). The middle column shows the location of KC shown using anti-F4/80 (red) in the top panel and the location of LSEC using mab anti-2.4G2 (red) in the bottom panel. The merged color image shows the colocalization of signal from \(\gamma\)-chain with signals from F-4/80 and 2.4G2 along with DIC and DAPI staining of nuclei. Scale bars, 10 \(\mu\)m. (B) Quantitative colocalization analysis was applied to all images represented in (A). The graph represents the percentage of signal from \(\gamma\)-chain Ab (green) colocalizing with signal from F4/80 or 2.4G2 (red) Ab and vice versa in the top panel of (A) and also for the bottom panel (mean ± SD). Using a random-effects linear regression model, the data from bars 1 and 3 and also between bar 2 and 4 were analyzed. *\( p < 0.001 \).

![Image 7](https://example.com/image7)

**FIGURE 7.** (A) Quantitative precipitin curve. The graph shows a quantitative precipitin curve plotting the protein concentrations of immune precipitates (pellets) from a series of tubes incubating increasing concentrations of Ag and a fixed concentration of Ab. The peak marks the POE. (B) Particle size measurement. Particle size distribution of SIC prepared at a 15-fold Ag excess was measured using DLS; the data obtained with Dynapro software are presented. The graph plots particle mean (± variance) diameter in nm on the x-axis, and the y-axis shows the percent distribution of total particles calculated using the relative amount of particles \([G(d)]\) at each size.
FIGURE 8. SIC are not cleared efficiently in the RIIb KO mouse strain. We infused via the tail vein radio-iodinated SIC prepared at 15× Ag excess of POE and then evaluated the clearance of SIC from peripheral blood by counting gamma-radiation in blood collected from the retro-orbital sinuses. The curve plots mean ± SD values of SIC in blood samples, after adjustment for dose/body weight (see Materials and Methods), over time from six WT (closed circle) and five RIIb KO (open circle) animals combining all data from two experiments done 3 mo apart. *p < 0.05 (statistically significant differences between strain values at each time point; Student t test).

The fraction of total RIIb on LSEC is astonishingly high, about three-quarters of the total RIIb in the body. Relatively small fractions of the total are noted in spleen and blood and bone marrow. It should be acknowledged that we have not sampled all organs, so our total value may be somewhat underestimated, but in fact we have sampled all of the major immune system organs and would conclude that the liver content is far in excess of that of any other organ.

The second major conclusion to be derived from our data is that RIIb of the liver sinusoidal endothelium is responsible for removing SIC from blood. This follows from our observation that the elimination rate of SIC from blood of the RIIb KO strain was severely diminished. Three elements of this conclusion require comment. First, we contend that the liver is the major organ removing SIC. This contention is supported by our observations 1) that the RIIb KO mouse is severely deficient in eliminating SIC, 2) that the majority of RIIb of the mouse is expressed in the liver, and 3) that in our hands 96% of cleared SIC is found in the liver by direct gamma-counting. Further, the literature supports this conclusion with abundant direct data showing liver clearance of the major fraction of blood-borne SIC (1, 3, 4, 15, 39, 40). The spleen, contrary to conventional wisdom, is only a minor participant in SIC elimination.

The second element of our second major conclusion is that RIIb is the operant FcγR responsible for liver clearance. This observation is remarkable because this receptor has been known almost exclusively as an inhibiting receptor, downregulating the immune complex-mediated effects of the other agonist receptors in its gene family. By contrast, here in our studies it functions as an agonist. Early work, however, indicated that it was indeed capable of agonist activity, specifically of pinocytic uptake of ligand; but these were in vitro experiments and not in vivo as ours are (41–43). We know of one other situation where this receptor appears to act alone, without an associated receptor, and that is to produce apoptosis when clustered on B cells (44). The mechanistic details of how RIIb acts on LSEC to eliminate immune complexes will require further study.

The third remarkable element of our second major conclusion is that a specialized endothelium, the LSEC, is the predominant cell responsible for SIC elimination in that 90% of liver RIIb is expressed on LSEC. The LSEC, according to the literature, is a vigorous scavenger of bloodstream detritus, expressing a variety of surface receptors for blood-borne material, displaying abundant coated vesicles and lysosomes appropriate for a disposal mechanism (45), and in our hands eliminating blood-borne virus with remarkable efficiency (12). Further, the uptake of immune complex by LSEC has been well documented by others (4, 13, 16, 46). We note parenthetically that FcγR-expressing endothelium has been found elsewhere only in the human placenta, and there, too, the isoform expressed is RIIb (28).

The absence of the small fraction (10%) of total liver RIIb expressed on KC cannot be responsible for the slow rate of immune complex elimination in the RIIb KO mouse. Rather, the absence of RIIb on KC would be predicted to render the KC more vigorously endocytic due to overactive agonistic FcγR unopposed by the inhibitory RIIb. The result would be a more rapid decay of blood-borne SIC in the KO strain compared with the WT strain, the exact reverse of what we found.

We find the decay kinetics of SIC in the RIIb KO strain noteworthy on two accounts. First is the striking decrease in the rate of elimination during the latter of the two biexponential elimination phases, from 5 to 60 min, indicating that RIIb is largely responsible for the removal of blood-borne SIC. Second is the observation that the 1-min point on the WT curve is >20% lower than the 1-min point of the KO curve (p = 0.002, Fig. 8) despite the strains not only being matched for age and sex but also adjusted for radioisotope dose and body weight; and, further, that the period from 1 to 5 min shows similar rates of decay in both strains. We interpret these kinetics to suggest that during the first minute after infusion, the LSEC RIIb become saturated with SIC that pinocytose into (likely) coated vesicles, disappearing from the LSEC surface such that for the next 5 min, no more RIIb-specific elimination occurs. Then, after 5 min, RIIb begin to reappear on the LSEC surface, either by re-expression of endocytosed receptor, suggested by others (16), or by biosynthesis; and an ongoing elimination of SIC ensues at a vigorous rate (half-life ~90 min). This supposition suggests a fruitful path of insight into the mechanism of elimination and warrants detailed investigation.

We call attention to old observations that the repetitive infusion of SIC, such as those used here in our study, causes serum sickness with characteristic microscopic findings of glomerulonephritis (6, 7, 47). We speculate that LSEC RIIb, mediating the elimination of SIC, may critically attenuate the manifestations of serum sickness. One would further anticipate that a strain of mouse lacking only LSEC RIIb may be more susceptible to serum sickness and other soluble immune complex-mediated diseases, whereas targeted elimination of RIIb in B cells and myeloid cells would not produce like effects. Likewise, animals harboring mutants of the RIIb gene that result in diminished or defective receptor expression may as well be more susceptible to immune complex-mediated diseases. Such mutants have been described although their analyses appear to have focused on B cell and myeloid RIIb expression, whereas the evaluation of RIIb expression on LSEC would be expected to reveal more telling results (20, 48, 49).

Summarizing for clarity, risking oversimplification, we would propose that SIC are constitutively eliminated by LSEC RIIb, but when this capacity is overwhelmed, the escaping immune complexes become large and lodge in end organs to result in disease. Alterations in the expression and function of RIIb would modify this course of events.
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Disclosures
The authors have no financial conflicts of interest.

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