

IgG hexamers initiate acute lung injury

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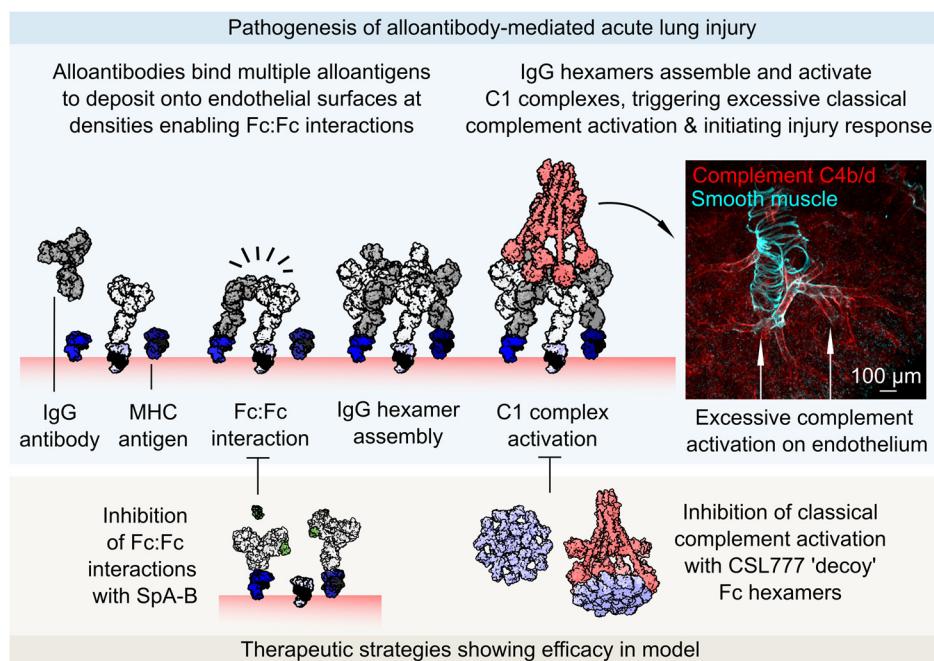
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Graphical abstract



1 **Abstract**

2 Antibodies can initiate lung injury in a variety of disease states such as autoimmunity, transfusion
3 reactions, or after organ transplantation, but the key factors determining in vivo pathogenicity of injury-
4 inducing antibodies are unclear. A previously overlooked step in complement activation by IgG
5 antibodies has been elucidated involving interactions between IgG Fc domains that enable assembly of
6 IgG hexamers, which can optimally activate the complement cascade. Here, we tested the in vivo
7 relevance of IgG hexamers in a complement-dependent alloantibody model of acute lung injury. We
8 used three approaches to block alloantibody hexamerization (antibody carbamylation, the K439E Fc
9 mutation, or treatment with domain B from Staphylococcal protein A), all of which reduced acute lung
10 injury. Conversely, Fc mutations promoting spontaneous hexamerization made a harmful alloantibody
11 into a more potent inducer of acute lung injury and rendered an innocuous alloantibody pathogenic.
12 Treatment with a recombinant Fc hexamer 'decoy' therapeutic protected mice from lung injury,
13 including in a model with transgenic human FCGR2A expression that exacerbated pathology. These
14 results indicate a direct in vivo role of IgG hexamerization in initiating acute lung injury and the potential
15 for therapeutics that inhibit or mimic hexamerization to treat antibody-mediated diseases.

16

17 **Brief summary**

18 IgG antibodies can form hexamers. This study shows that hexamer assembly is an important event
19 determining the ability of IgG to trigger acute lung injury.

20 **Introduction**

21 Antibodies and the complement cascade mediate protective immunity but can both become misdirected
22 to cause harm in autoimmune and alloimmune diseases. Some antibodies can direct activation of the
23 complement cascade at their targets, an event associated with severe pathology in disease states
24 including several forms of transfusion reactions (1, 2), immune rejection after solid organ
25 transplantation (3), and complications of pregnancy (4). Complement-activating alloantibodies are
26 known mediators of transfusion-related acute lung injury (TRALI) (5, 6), a leading cause of transfusion-
27 related deaths (7), and are linked to particularly poor outcomes following solid organ transplantation (3,
28 8). Complement activation by autoreactive antibodies also contributes to pathogenesis of forms of
29 autoimmune hemolytic anemia (9), small vessel vasculitis (10), and neurological autoimmune disease
30 (11).

31 Immunoglobulin G (IgG) antibodies are the most prevalent type of antibody in circulation and
32 complement-activating alloantibodies are frequently IgG class. IgG antibodies achieve complement
33 activation through recruiting and activating C1 complexes, each of which contain six Fc-binding
34 domains (12, 13). A theory for how IgG achieves C1 complex activation involves groups of six IgG
35 antibodies interacting through their Fc domains to form IgG hexamers (14). This theory recently gained
36 experimental support from direct imaging of IgG1 and IgG3 hexamer assembly on antigenic liposomes
37 (15, 16), with in vitro studies connecting IgG hexamerization to increased complement deposition on
38 target surfaces (15, 17, 18). However, it is currently unclear whether IgG hexamer assembly is
39 important in vivo in the pathogenesis of complement-dependent forms of alloantibody-mediated
40 disease.

41 Here, we report testing of interventions that exploit IgG hexamerization in a mouse model of acute lung
42 injury driven by alloantibody deposition in the pulmonary microvasculature, a process that drives
43 pathology in forms of both TRALI and antibody-mediated rejection (AbMR) of lung transplants (6). Our
44 results identify key molecular events driving alloantibody-mediated pathophysiology in vivo. We also
45 demonstrate preclinical efficacy of new therapeutic approaches that prevent pathology of complement-
46 dependent organ damage caused by alloantibodies, serving as a rationale to pursue translational
47 studies in human alloantibody driven disease.

48 **Results**

49 Alloantibodies are prevalent but not always harmful, so determining whether alloantibodies are clinically
50 significant is a frequent conundrum in transfusion and transplantation medicine. Reflecting this clinical
51 challenge, of the many mouse monoclonal alloantibodies targeting major histocompatibility complex
52 (MHC) class I antigens, only clone 34-1-2S triggers acute lung injury when microgram quantities are
53 intravenously injected into mice (19, 20). In addition, only mice expressing the H-2^d MHC class I
54 haplotype are known to be susceptible to injury caused by the 34-1-2S antibody (5, 6). Curiously, the
55 34-1-2S antibody does not readily cause injury in H-2^b mice, including the widely used C57BL/6 (B6)
56 strain, despite the fact that it binds to MHC class I antigens expressed by H-2^b mice (5, 6). We aimed to
57 measure the affinity of 34-1-2S for a range of MHC class I antigens to improve our understanding of the
58 factors determining the ability of antibodies to cause injury in both this widely used model and more
59 generally in antibody-mediated disease states.

60 We measured the binding affinity of 34-1-2S antibody to each of the classical MHC class I antigens
61 present on injury-resistant H-2^b B6 mice and injury-susceptible H-2^d mice (**Fig. 1A**). Of the three MHC
62 antigens in the H-2 locus (K, D, and L), B6 mice only express K^b and D^b, and we detected binding of
63 34-1-2S to K^b but not D^b. In contrast, we detected binding of 34-1-2S to all three MHC class I antigens
64 from H-2^d mice, with high affinity binding to K^d and D^d, and weak binding to L^d (**Fig. 1B**). Other MHC
65 class I antibodies (clones AF6-88.5.5.3, 20-8-4S, SF1.1.10, 30-5-7S and 34-5-8S), which do not readily
66 induce injury (5) each bound to only one MHC class I antigen from each MHC type (**Fig. S1**).

67 Together, the above findings led us to the hypothesis that the ability of 34-1-2S to induce lung injury in
68 H-2^d mice is a function of increased density of bound antibody in H-2^d animals resulting from 34-1-2S
69 simultaneously binding K^d, D^d, and possibly L^d. This hypothesis was tested by injecting 34-1-2S
70 antibody into B6.ConK^d-on mice, which express K^d but do not express D^d or L^d (21). B6.H2^d mice
71 expressing the full complement of MHC class I antigens recognized by 34-1-2S (K^d, D^d, and L^d) were
72 used as background-matched positive controls for susceptibility to injury (**Fig. 1C**). In contrast to
73 B6.H2^d mice, B6.ConK^d-on mice did not develop lung injury (**Fig. 1D, E**). These data are consistent with
74 34-1-2S antibody causing injury in H2^d mice through high affinity binding to multiple MHC class I
75 antigens.

76 Engagement of multiple antigens can permit high density antibody deposition, an event associated with
77 classical complement activation. Complement activation has been implicated in pathogenesis of acute
78 lung injury caused by 34-1-2S antibody, but previous studies have not determined whether injury in this
79 model is directly triggered by antibody-mediated complement activation via the classical pathway (5, 6).

80 To test whether 34-1-2S-induced injury requires classical complement activation, we bred mice
81 expressing the *H2^d* susceptibility locus with mice lacking C1qa (22), a protein that is necessary for
82 classical complement activation as it is one of the three proteins which make up each of the six Fc-
83 binding C1q subcomponents in each C1 complex (**Fig. 2A**).

84 Relative to B6.*H2^d:C1qa^{+/+}* littermate controls, C1qa-deficient B6.*H2^d:C1qa^{-/-}* mice were resistant to
85 alloantibody-mediated acute lung injury and mortality (**Fig. 2B-D**). Mice lacking C1qa were also
86 protected from deposition of complement component C3 split products on the endothelium of
87 pulmonary capillaries (**Fig. 2E**). Staining for C1qa in lungs confirmed absence of C1qa protein in
88 knockout mice, with intense C1 complex deposition seen around pulmonary arteriolar endothelial cells
89 in *C1qa*-expressing mice injected with 34-1-2S antibody (**Fig. 2F**).

90 To identify the microanatomical site of classical complement activation, we stained lungs of mice
91 injected with 34-1-2S for the complement split products C4b and C4d, which form covalent bonds with
92 proteins at sites of classical complement activation. We observed strong positivity for C4b/d highlighting
93 the endothelium of medium and small-sized pulmonary arterioles in B6.*H2^d:C1qa^{+/+}* mice injected with
94 34-1-2S, but not in B6.*H2^d:C1qa^{-/-}* mice (**Fig. 2G** and **Movie S1**). Together, these results indicate that
95 34-1-2S causes acute lung injury because this antibody is deposited onto the pulmonary arteriolar
96 endothelium at densities sufficient to trigger excessive classical complement activation directed at the
97 walls of these blood vessels.

98 Dense binding to membrane-expressed antigens would be expected to facilitate IgG Fc:Fc interactions
99 and IgG hexamer assembly. IgG hexamers are potent activators of C1 complexes in vitro (15), and are
100 further implicated in classical complement activation by models for C1 complex activation involving
101 shifting of its six Fc-binding C1q subcomponents into a regular hexagonal configuration (**Fig. 2A, 3A**)
102 (12, 13). We therefore hypothesized that 34-1-2S assembles into hexamers on the pulmonary
103 endothelial surface of susceptible mice to trigger complement-dependent acute lung injury.

104 Imaging methods cannot currently resolve IgG hexamers *in vivo*, but recent studies have developed
105 methods for inhibiting IgG hexamerization. One approach to impair IgG hexamer assembly is to
106 carbamylate antibodies, converting lysine residues to homocitrullines to alter charge densities in IgG Fc
107 regions, inhibiting Fc:Fc interactions and IgG hexamer assembly (**Fig. 3B**) (23). Mice treated with
108 carbamylated 34-1-2S showed greatly reduced acute lung injury responses compared to littermate
109 controls treated with non-carbamylated 34-1-2S (**Fig. 3C, D**). Carbamylated 34-1-2S was deposited in
110 lungs but, in contrast to unchanged 34-1-2S, did not induce complement C3b/d deposition in the
111 pulmonary microvasculature (**Fig. 3E**)

112 Lysine residues are present on regions of IgG outside of the Fc:Fc interaction interface (illustrated in
113 **Fig. 3B**), so we pursued a more targeted strategy for inhibition of IgG hexamer assembly. We
114 determined the sequence of both heavy and light chain complementary-determining regions and
115 engineered a chimeric antibody with the Fab domain of 34-1-2S fused in frame to human IgG1 (hlgG1-
116 34-1-2S). To test whether hlgG1-34-1-2S causes injury through hexamerization, we also expressed this
117 antibody with an Fc point mutation that inhibits Fc:Fc interactions required for IgG hexamer assembly
118 (K439E, **Fig. 3F**) (15). hlgG1-34-1-2S caused acute lung injury that was reduced by the K439E
119 mutation (**Fig. 3G, H**), as was complement C4b/d deposition in lungs (**Fig 3I**), lending further support to
120 a role for Fc:Fc interactions and hexamerization in the pathogenesis of this disease model.

121 We also tested a strategy for pharmacologic inhibition of Fc:Fc interactions by mixing hlgG1-34-1-2S
122 with recombinant B domains from *Staphylococcus aureus* protein A (SpA-B), which bind to IgG
123 antibodies near to Fc:Fc interaction sites and inhibit hexamer assembly and complement activation by
124 antibodies targeting bacterial antigens (**Fig. 3J**) (17, 24). We hypothesized that these properties of
125 SpA-B, which likely evolved as part of an immune evasion strategy, might be harnessed to prevent
126 hlgG1-34-1-2S from causing acute lung injury. Adding SpA-B to hlgG1-34-1-2S reduced its ability to
127 both induce acute lung injury (**Fig. 3K, L**) and cause complement C4b/d deposition within pulmonary
128 arterioles (**Fig. 3M**). These findings provide a third line of evidence that Fc:Fc interactions leading to
129 hexamer assembly are important for the injury response caused by this alloantibody.

130 Turning to hexamer gain of function experiments, the introduction of three mutations into the Fc domain
131 of hlgG1 (RGY mutations: E345R, E430G, S440Y) resulted in antibodies capable of off-target hexamer
132 assembly as well as increased on-target hexamerization (15) (**Fig. 4A**). We hypothesized that RGY-
133 mutated 34-1-2S (RGY-hlgG1-34-1-2S) would have enhanced ability to cause acute lung injury due to
134 increased IgG hexamer formation. We produced RGY-hlgG1-34-1-2S and confirmed its ability to
135 spontaneously assemble into hexamers in solution (**Fig. 4B**). Consistent with a role for alloantibody
136 hexamerization in driving injury, RGY-hlgG1-34-1-2S showed increased potency in triggering acute
137 lung injury relative to hlgG1-34-1-2S (**Fig. 4C, D**), and inducing complement C4b/d deposition in lungs
138 (**Fig. 4E**). A novel chimeric hlgG1 antibody binding a single MHC class I antigen (hlgG1-Kd1, targeting
139 K^d), did not provoke injury when injected into B6.H2^d mice (**Fig. 4F, G**), consistent with binding to
140 multiple antigens being a requirement for alloantibody-mediated acute lung injury. However,
141 introduction of the RGY mutations promoting hexamerization into this innocuous antibody resulted in a
142 modified version (RGY-hlgG1-Kd1) that was able to provoke increases in lung vascular permeability
143 and edema (**Fig. 4F & G**). Promoting IgG hexamer assembly can therefore increase in vivo
144 pathogenicity of alloantibodies.

145 Another approach for therapeutic exploitation of IgG hexamerization involves use of Fc hexamers as
146 'decoy' treatments. These therapeutic candidates are under investigation as recombinant alternatives to
147 plasma-derived intravenous or subcutaneous immunoglobulin (IVIg or SC Ig) treatments that are used in
148 management of autoimmune and alloimmune diseases (25). We hypothesized that due to its ability to
149 inhibit classical complement activation (11, 26), the Fc hexamer 'decoy' treatment CSL777 (previously
150 Fc- μ TP-L309C) would be effective in preventing alloantibody-mediated acute lung injury.

151 We randomized mice to receive either CSL777, SC Ig (IgPro20, a human plasma-derived
152 immunoglobulin product which is currently used to treat antibody-mediated diseases), or vehicle
153 controls prior to injection with 34-1-2S (**Fig. 5A, B**). Treatment with CSL777 protected mice from
154 developing 34-1-2S-induced lung vascular permeability and pulmonary edema responses at all doses
155 tested, whereas treatment with SC Ig only had a partial effect on alloantibody-induced acute lung injury
156 responses (**Fig. 5C-F**). CSL777 treated mice lacked alloantibody-mediated deposition of complement
157 C4 split products on pulmonary arterioles, whereas arteriolar endothelial C4b/d deposition was still
158 observed in lungs of SC Ig-treated mice after 34-1-2S antibody injections (**Fig. 5G, H**). Recombinant Fc
159 hexamer therapeutics such as CSL777 might therefore be useful for prevention or treatment of
160 complement-dependent forms of alloantibody-mediated organ injury.

161 Unlike humans, mice do not express the Fc γ receptor FCGR2A (Fc γ RIIA, CD32A), negatively
162 impacting the predictive value of mouse models for studying human antibody-mediated diseases (27,
163 28). To test whether our previous findings held up in a system involving FCGR2A-driven pathology, we
164 crossed existing mouse lines to generate 34-1-2S-mediated injury-susceptible H-2 d mice expressing a
165 human FCGR2A (hFCGR2A) transgene (B6.H2 d :hFCGR2A $^{Tg/0}$). Mice expressing hFCGR2A developed
166 similar levels of lung injury relative to littermates without hFCGR2A expression but displayed a survival
167 disadvantage (**Fig. 6A-C**), and increased sequestration of platelets in the pulmonary microvasculature
168 (**Fig. 6D,E**). Classical complement activation was still critical for pathogenesis in the presence of
169 hFCGR2A, as knockout of C1qa protected mice expressing hFCGR2A from lung injury and mortality
170 (**Fig. 6F-H**). B6.H2 d :hFCGR2A $^{Tg/0}$ mice treated with CSL777 were protected from 34-1-2S-mediated
171 injury and showed no mortality responses (**Fig. 6I-K**). These results provide evidence that in a
172 humanized antibody-mediated acute lung injury model involving pathology driven by hFCGR2A,
173 classical complement activation is still a critical event in pathogenesis that can be targeted by
174 therapeutics.

175 **Discussion**

176 This study advances our understanding of immunology in three areas. Our work elucidates the
177 molecular determinants of susceptibility in a widely used inflammation model. To our knowledge, our
178 experiments represent the first *in vivo* evidence that alloantibody hexamerization is important for
179 pathophysiology. In addition, we show that two experimental therapeutic approaches that target
180 antibody hexamerization can prevent alloantibody-mediated organ injury.

181 The findings presented in this study allow us to explain a long-standing mystery of great mechanistic
182 importance in a widely used model of immune-mediated organ injury; in particular, why 34-1-2S
183 antibody injections (but not injections of other anti-MHC class I antibodies) cause such striking
184 pathophysiology in mice carrying the H-2^d but not H-2^b MHC haplotype (6, 19, 20, 28–38). We posit that
185 high affinity binding to multiple MHC class I antigens on the pulmonary endothelium of mice with the H-
186 2^d haplotype facilitates sufficiently dense alloantibody deposition for IgG hexamer assembly to occur.
187 These IgG hexamers then direct classical complement activation onto the pulmonary endothelial
188 surface, initiating the excessive leukocyte and platelet responses that have been reported in previous
189 studies to cause acute lung injury in this model (6, 19, 20). Furthermore, we were able to render an
190 innocuous antibody specific for a single MHC class I antigen into a pathogenic antibody by introducing
191 mutations promoting increased on and off-target hexamerization. An additional implication of our
192 findings is that both lymphocyte crossmatch and single antigen bead assays for detecting complement-
193 fixing antibodies may lack sensitivity for detecting antibodies that activate complement *in vivo*. The
194 mobility, density and diversity of antigens on the lymphocytes or solid-phase beads used in these
195 assays does not exactly resemble those on endothelial cell surfaces targeted by donor-specific
196 antibodies *in vivo*, and our results indicate that each of these factors determines complement-fixing
197 capability of antibodies. Conversely, antigen density on beads exceeding that found *in vivo* may give
198 false positive findings of complement-fixing antibody responses.

199 By demonstrating that IgG hexamers are important in pathophysiology and represent therapeutic
200 targets *in vivo*, our work builds on *in vitro* studies implicating antibody hexamerization in complement
201 activation by antibodies targeting antigens on liposomes, lymphoma cells or bacterial membranes (15–
202 18, 23, 39). Further clinical translation will require determination of the importance of IgG hexamers in
203 more complex models of diseases involving complement-activating alloantibodies (e.g. AbMR, TRALI,
204 hemolytic transfusion reactions, and hemolytic disease of the fetus and newborn) or autoantibodies
205 (e.g. warm autoimmune hemolytic anemia, antiphospholipid syndrome, myasthenia gravis and
206 neuromyelitis optica). As SpA-B does not inhibit IgG3-mediated complement activation but IgG3 can

207 assemble into hexamers (15, 16), it will also be important to develop strategies to inhibit IgG3
208 hexamerization to examine the therapeutic potential of targeting hexamers formed by IgG3 antibodies.
209 Our results also provide new insights into the modes of action of past, present, and potential future
210 therapeutics. Full length staphylococcal protein A (SpA) has been used as a therapeutic in the form of a
211 now-discontinued extracorporeal immunoabsorption product (Prosorba column). Efficacy of SpA
212 immunoabsorption has been observed in patients with symptoms unchanged by plasma exchange, an
213 effect ascribed to leakage of SpA from columns into the bloodstreams of patients resulting in B cell
214 depletion caused by the action of SpA as a B cell receptor super agonist (40). Purified SpA infusions
215 (PRTX-100) were subsequently studied in early-stage clinical trials before abandonment for financial
216 reasons (41). Our results suggest that there may be settings where therapeutics based on the SpA-B
217 subdomain of SpA have efficacy through preventing complement activation without risk of adverse
218 effects related to immune complex formation and B cell super agonism caused by immunoglobulin
219 polyvalency of full-length SpA. Donor-derived immunoglobulin products (e.g. IVIg and SCIG) are
220 currently used to treat antibody-mediated disease flares. Our observation that SCIG reduces injury
221 responses but does not prevent classical complement activation in vivo is concordant with previous
222 studies concluding that immunoglobulin therapeutics act on downstream mediators in vitro and in vivo
223 (33, 42). CSL777 is an attractive potential future therapeutic for treatment of alloantibody-mediated
224 diseases as it showed efficacy in our models, and had a mode of action that would be anticipated to
225 prevent complement activation by both IgM and IgG antibodies as well as pathophysiology resulting
226 from Fc γ receptors (11, 25, 26). CSL777 also lacks issues with use of donor-derived products related to
227 sourcing, purification and concentration for injections (25, 43).
228 In conclusion, this study provides evidence that IgG hexamers can be important triggers of
229 complement-dependent pathophysiology in vivo. Our preclinical studies support further investigation of
230 IgG hexamerization inhibitors and IgG hexamer 'decoy' therapeutics for use in preventing disease
231 states caused by antibodies and complement activation.

232 **Materials and methods**

233 *Animals*

234 B6.C-H2^d/bByJ (B6.H2^d) mice (Cat# 000359) (44) and B6(Cg)-C1q $\alpha^{tm1d(EUCOMM)Wtsi}$ /TennJ (C1q $\alpha^{-/-}$) mice
235 (Cat# 031675) (22) originated from the Jackson Laboratory. B6-background mice were bred with
236 B6.H2^d mice and progeny were crossed to produce mice with homozygous expression of H2^d MHC
237 antigens for use in experiments (6). B6.ConK^d-on and B6.Tg(CD2-Tcra,-Tcrb)75Bucy (TCR75) mice
238 were provided by J. Zimring (21, 45). BALB/c mice were from Charles River Laboratories (Cat# 028).
239 B6;SJL-Tg(FCGR2A)11Mkz/J mice (expressing human FCGR2A isoform R131, Jackson Laboratory
240 Cat# 003542) (27) were backcrossed to B6 congenicity (46). Male mice were used as female mice are
241 not susceptible to 34-1-2S-mediated injury (5). Mice were studied at 8-16 weeks of age after
242 maintenance in the UCSF specific pathogen-free animal facility for at least 2 weeks. Procedures
243 received ethical approval from the UCSF IACUC committee.

244 *Surface plasmon resonance*

245 Binding affinities were determined by injecting serial dilutions (0.5-200 nM) of MHC class I monomers
246 (MBL International Cat#: TB-5001-M (K^b presenting SIINFEKL); TB-5008-M (D^b presenting
247 RAHYNIVTF); TB-M552-M (K^d presenting VYLKTNVFL); TB-M536-M (D^d presenting IGPGRAFYA);
248 TB-M521-M (L^d presenting SPSYVYHQF)) over monoclonal antibodies bound via amine coupling to
249 SensEye G Easy2Spot sensors (Ssens Cat# 1-09-04-006), assayed in triplicate with an IBIS MX96
250 SPR imager.

251 *Alloantibody-mediated acute lung injury model*

252 As previously described, mice were given i.p. injections of LPS (Sigma Aldrich Cat# L2880, 0.1 mg/kg)
253 for 'priming' needed to render barrier-housed mice responsive to antibody injections (19). At 24 hours
254 after LPS priming, mice were anesthetized (0.6 mg/kg ketamine + 0.1 mg/kg xylazine i.p.) and the
255 indicated antibody treatments were injected into the jugular vein over the course of 1 minute (at 1
256 mg/kg unless otherwise stated). Antibodies were from BioXCell (mlgG2a isotype control Cat# BE0085,
257 34-1-2S Cat# BE0180, hIgG1 isotype control Cat# BE0297) or newly produced (described below). Lung
258 vascular permeability was measured by giving each mouse 0.01 KBq of ¹³¹I-conjugated albumin (Iso-
259 Tex Diagnostics, NDC:50914-7731) together with i.v. antibody injections, collecting lungs and blood
260 samples 2 hours later or at cessation of breathing for radioactivity measurements used to quantify
261 volume of extravasated plasma in lungs (lung vascular permeability). Wet-dry weight ratios of lungs and
262 blood were used to calculate excess lung water volumes (6).

263 *Immunofluorescence imaging*

264 Cryosections were made at 200 μ m or 400 μ m thickness from lungs fixed by inflation with and
265 immersion in 1% formaldehyde in PBS, as previously described (6). Sections were incubated overnight
266 with antibodies targeting C3b/d (Novus Cat# NB200-540), C1qa (Abcam Cat# ab182451), C4b/d
267 (Novus Cat# NB200-541), Scgb1a1 (Sigma Aldrich Cat# ABS1673) or CD41 (Biolegend Cat# 133939)
268 together with a FITC-conjugated antibody raised against Acta2 (Sigma-Aldrich Cat# F3777) were
269 incubated overnight at 1:500 with 5% normal donkey serum, 0.1% bovine serum albumin and 0.3%
270 triton X-100 in phosphate-buffered saline (PBS). After washing, Cy3 or Alexa Fluor 647-conjugated
271 cross-adsorbed polyclonal secondary antibodies targeting rat, rabbit, goat and/or mouse IgG (Jackson
272 Immunoresearch Cat# 712-165-153, Cat# 711-165-152, Cat# 705-605-147 and/or Cat# 115-605-206)
273 were incubated with sections at 1:500 in PBS + 0.3% triton X-100 overnight. After additional washes,
274 sections were either mounted in Vectashield (Vector Laboratories Cat# H-1700) for standard confocal
275 imaging on a Nikon A1r microscope, or cleared after staining using the EZ clear protocol (47) for 3D
276 imaging with a Nikon AZ100M confocal microscope.

277 *Antibody carbamylation*

278 Carbamylation of 34-1-2S was achieved by incubating 1 mg of antibody in PBS + 0.1 M KOCN for 1
279 hour at 37°C before buffer exchange back into PBS (23). Control 34-1-2S was subjected to the same
280 process with omission of KOCN.

281 *Antibody sequencing and engineering*

282 The 34-1-2S hybridoma was purchased from ATCC (Cat# HB-79). To generate the Kd1 hybridoma, B6
283 mice were injected i.v. with 3×10^6 CD4+ T cells from TCR75 mice transgenic for a T cell receptor (TCR)
284 specific for a peptide from K^d presented by IA^b, and i.p. with 5×10^6 Con-K^d-on splenocytes, resulting in
285 an extreme B cell antibody response directed at an immunodominant peptide from K^d, the only
286 mismatched antigen between donor and recipient. Three days after a boost with an additional i.p.
287 injection with Con-K^d-on splenocytes, splenocytes from the sensitized recipient were fused with a
288 myeloma cell line as previously described (48), and monoclonal antibodies specific for K^d were
289 identified using Con-K^d-on splenocytes as targets (21).

290 Monoclonal antibody aliquots were digested with either peptidyl-Asp metalloendopeptidase,
291 chymotrypsin, elastase, trypsin, or pepsin enzymes. Peptides were then assayed using liquid
292 chromatography coupled to tandem mass spectrometry for sequencing of variable fragments
293 (Bioinformatics Solutions Inc.). Amino acid sequences determined were, for 34-1-2S:

294 EVQLQQSGAEFVRPGASVQLSCTASGFNLKDDYMFVVKQRPEQGLEWIGWIAPDNGDTEYASKFQG
295 KATITADTSSNTAYVQLSSLTSEDTAVYYCTTWGYYSYVNYWGQGTTLVSS (heavy chain variable
296 region) and:
297 DIQMTQSPSSLSASLGERVSLTCRASQDIGSNLNWLQQEPDGTIKRLIYATYSLDSGVPKRFSGSRSGS
298 DYSLTISSEDFVDYYCLQYASSPYTFGGGTKEIK (light chain variable region); and for Kd1:
299 EVLLVESGGDLVKPGGSLKLSCAASGFTFRYAMSWVRQTPEKRLIEWVATIGDDGSYTFYPDNVKGR
300 FTISRDNAKNNLYLQMRHLKSEDTAIYFCARDGLFAYWGQGTLTVSA (heavy chain variable region)
301 and:
302 DIQMTQSPSSLSASLGGKVITCKASQDIKKNIAWYQYKPGKGPRLLIHYTSTLQPGISSLRFSGSGSGR
303 DYSFSISNLEPEDIATYYCLQYDSLLYTFGGGTKEIK (light chain variable region). Correct
304 identification of variable domains was confirmed by performing sequencing of the products of 5' rapid
305 amplification of cDNA ends (5'-RACE) for heavy and light chain mRNA using RNA isolated from
306 hybridomas. The isolated 5'-RACE amplicons contained open reading frames that encoded the above
307 peptides sequenced by mass spectrometry. These sequences were codon-optimized and antibodies
308 were expressed as chimeric hIgG1 with or without Fc point mutations in a HEK293 cell system and
309 purified using protein A and buffer exchange (Absolute Antibody).

310 *Pharmacologic treatments*

311 Recombinant subdomain B from *Staphylococcus aureus* (SpA-B, amino acid sequence:
312 HHHHHADNKFNKEQQNAFYEILHLPNLNEEQRNGFIQSLKDDPSQSANLLAEAKKLNDQAPK, His-
313 tag added for purification) was produced in an *E. coli* system by Genscript and supplied in protein
314 storage buffer (50 mM Tris-HCl, 150 mM NaCl, 10% Glycerol, pH 8.0). At 1-2 hours before i.v. injection,
315 SpA-B (3 mg/kg) or vehicle were mixed with hIgG1-34-1-2S resulting in a 30% vehicle, 70% PBS
316 mixture.

317 Trial formulations of CSL777 (in PBS vehicle), as well as clinical-grade IgPro20 (Hizentra™) and the
318 proprietary vehicle for IgPro20 were provided by CSL Behring. Mice were given i.p. injections of
319 CSL777, IgPro20 or relevant vehicle 1-2 hours before i.v. injections of antibodies at stated doses.

320 *Negative stain electron microscopy*

321 Antibody samples were diluted to 0.01 mg/ml in 25 mM HEPES, 100 mM NaCl and added to carbon-
322 coated grids (TedPella Cat# 01702-F, manually coated with 20 nm carbon using a Cressington 208
323 Sputter Coater). Sample-coated grids were stained using 0.75% uranyl formate and imaged on an FEI
324 Tecnai T12 transmission electron microscope.

325 *Experimental design and analysis*

326 Within-cage randomization was used for group allocations in studies testing exogenous treatments.
327 Littermate controls from heterozygous crosses were used to test the effect of *C1qa* knockout. Congenic
328 animals housed in the same room were used to study haplotype effects. Handlers were blinded to
329 group allocations during experiments. Group numbers (n) and analysis approaches were
330 predetermined before initiation of experiments. Statistical tests used on each dataset are described in
331 figure legends.

332 *Software*

333 GraphPad Prism was used for graphing and statistical analysis. UCSF ChimeraX was used for
334 molecular graphics (49). Imaris was used to render fluorescence micrographs and ImageJ was used to
335 process electron microscopy data.

336 **Acknowledgements**

337 Imaging studies were possible thanks to the UCSF Biological Imaging Development Colab, and UCSF
338 Electron Microscopy core. Sandro Prato (CSL Innovation Pty Ltd, Parkville, Victoria, Australia) provided
339 advice on in vivo administration of CSL777 and SC Ig.

340 **Funding**

341 National Institutes of Health grant R01 HL138673 (JCZ and MRL)
342 National Institutes of Health grant R35 HL161241 (MRL)
343 Association for Advancement of Blood and Biotherapies (AABB) Foundation (formerly National Blood
344 Foundation) Early Career Scientific Research Grant (SJC)
345 Canadian Institutes of Health Research (ÉB)

346 **Authorship**

347 Conceptualization: SJC, JCZ, MRL
348 Methodology: SJC, MRL, AEHB, GV, ÉB, RS, JCZ
349 Investigation: SJC, YS, JJT, NK, DPB, AEHB
350 Funding acquisition: SJC, JCZ, MRL
351 Writing – original draft: SJC, MRL
352 Writing – review & editing: SJC, YS, JJT, NK, DPB, AEHB, GV, ÉB, RS, JCZ, MRL
353 Conflict of interest statement: RS is an employee of CSL Behring AG, Switzerland. The authors have
354 no additional conflicts of interest.

355 **Data and materials availability**

356 All data are available in the main text or the supplementary materials.

357 **Supplementary materials**

358 Supplementary materials and methods.
359 Supplementary Figures:
360 **Fig. S1. Binding of MHC class I monoclonal antibodies to MHC class I monomers.**
361 **Movie S1. Complement C4 split product deposition in the pulmonary vasculature.**

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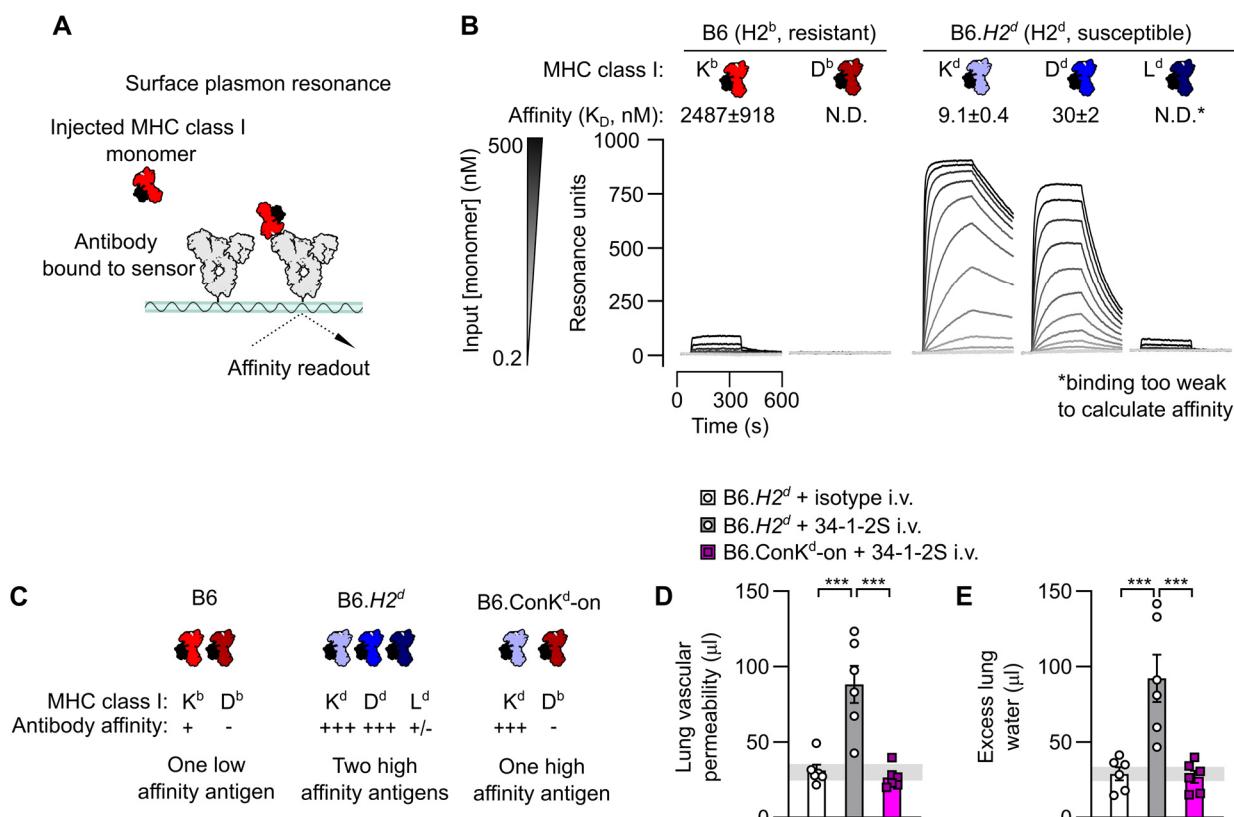
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468



469

470 **Fig. 1. The 34-1-2S alloantibody binds to multiple MHC class I antigens to trigger acute lung injury.**

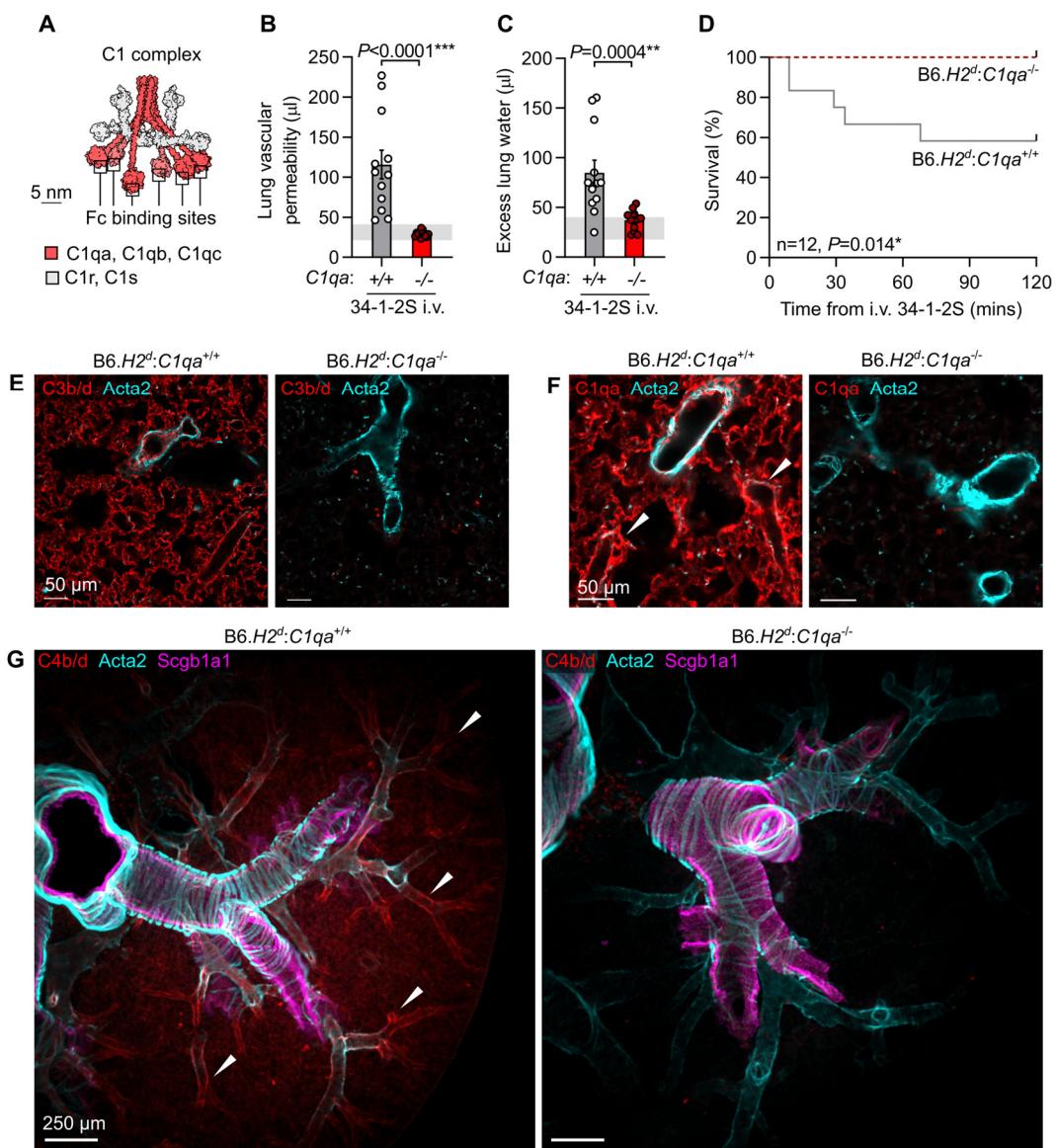
471 **A.** Schematic showing approach for measuring affinity of the MHC class I alloantibody 34-1-2S for MHC class I monomers
472 using surface plasmon resonance (SPR).

473 **B.** SPR sensorgrams showing detection of binding of 34-1-2S to the K^b MHC class I antigen from H2^b mice and the K^d, D^d and
474 L^d antigens from mice with the H2^d haplotype.

475 **C.** Classical MHC class I antigens present in B6, B6.H2^d and B6.Con-K^d-on mice with summary of results from **B**.

476 **D.** Lung vascular permeability and **E.** excess lung water measurements from LPS-primed B6.H2^d mice given intravenous (i.v.)
477 doses of either 34-1-2S or isotype control, versus B6.Con-K^d-on mice given i.v. 34-1-2S.

478 Depictions of IgG and MHC class I in **A-C** are based on protein data bank (PDB) entries 1HZH and 1RK1. **B, D & E** show
479 means ± standard errors. Statistical tests used on **D & F** were ordinary one-way ANOVA with Dunnett's test for differences
480 relative to B6.H2^d + 34-1-2S i.v. group, with data log₁₀-transformed prior to analysis, *** = P<0.0001.



481

482 **Fig. 2. Classical complement activation on the pulmonary endothelium initiates 34-1-2S-induced acute lung injury.**

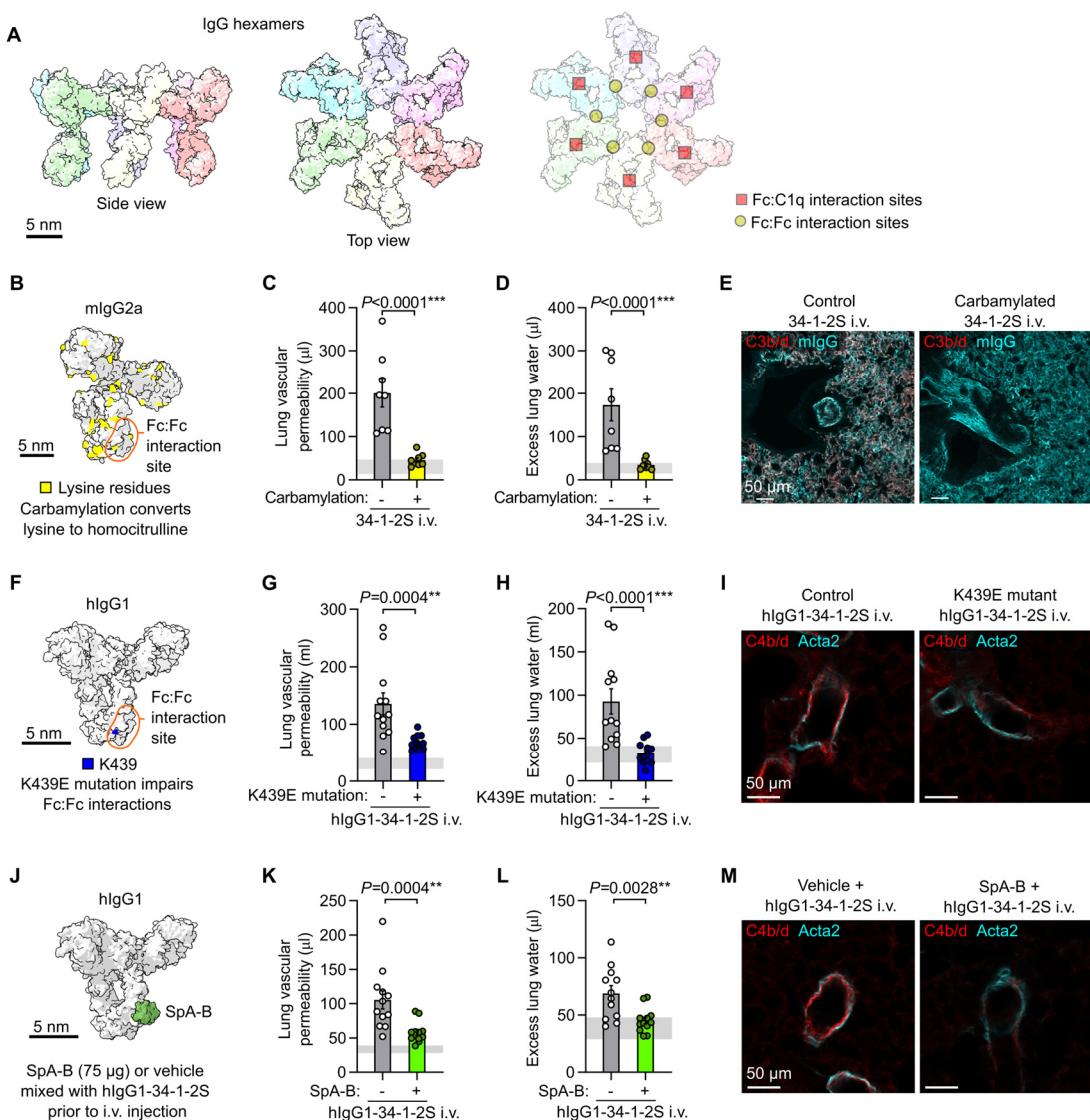
483 **A.** Molecular model of C1 complex based on small angle scattering database entry SASDB38 (12).

484 **B.** Lung vascular permeability and **C.** excess lung water measurements from LPS-primed B6.H2d:C1qa-/- mice and B6.H2d:C1qa^{+/+} littermates given i.v. 34-1-2S at 1 mg/kg. Horizontal gray lines are standard deviations of values from 'no injury' controls (B6.H2d mice given LPS i.p. + mlgG2a isotype control i.v.)

486 **D.** Survival of LPS-primed B6.H2d:C1qa-/- mice and B6.H2d:C1qa^{+/+} littermates given i.v. 34-1-2S at 4.5 mg/kg.

487 **E.** Immunofluorescence staining for complement C3b/d, **F.** C1qa or **G.** C4b/d (red) as well as Acta2 (α -smooth muscle actin, cyan) and, in **G.**, Scgb1a1 (club cell secretory protein, magenta) in lung sections from LPS-primed B6.H2d:C1qa-/- mice and B6.H2d:C1qa^{+/+} mice fixed 5 minutes after i.v. 34-1-2S at 1 mg/kg. Images in **G.** are maximum intensity projections sampling 240 μ m from a cleared thick section. White arrowheads point to arterioles positive for complement components.

492 **B & C** show means \pm standard errors. *P*-values are from unpaired two-tailed t-tests on log₁₀-transformed data (**B & C**) or log-rank test (**D**), with group n=12.



494

495 **Fig. 3. Inhibiting IgG hexamer assembly reduces 34-1-2S-induced acute lung injury responses.**

496 **A.** Molecular models of IgG hexamers based on PDB entry 1HZH, showing Fc:Fc and Fc:C1q interaction sites.

497 **B.** Molecular model showing lysine residues in mouse IgG2a (mlgG2a), PDB entry 1IGT.

498 **C.** Lung vascular permeability, **D.** excess lung water measurements and **E.** lung complement C3b/d and mlgG immunostains from LPS-primed BALB/c mice after i.v. injection of carbamylated or control 34-1-2S.

499 **F.** Molecular model showing location of Fc domain lysine 439 (K439) in human IgG1 (hlgG1), PDB entry 1HZH.

500 **G.** Lung vascular permeability, **H.** excess lung water measurements and **I.** lung complement C4b/d and Acta2 immunostains from LPS-primed B6.H2d mice after i.v. injection with hlgG1-34-1-2S or hlgG1-34-1-2S with K439E mutation.

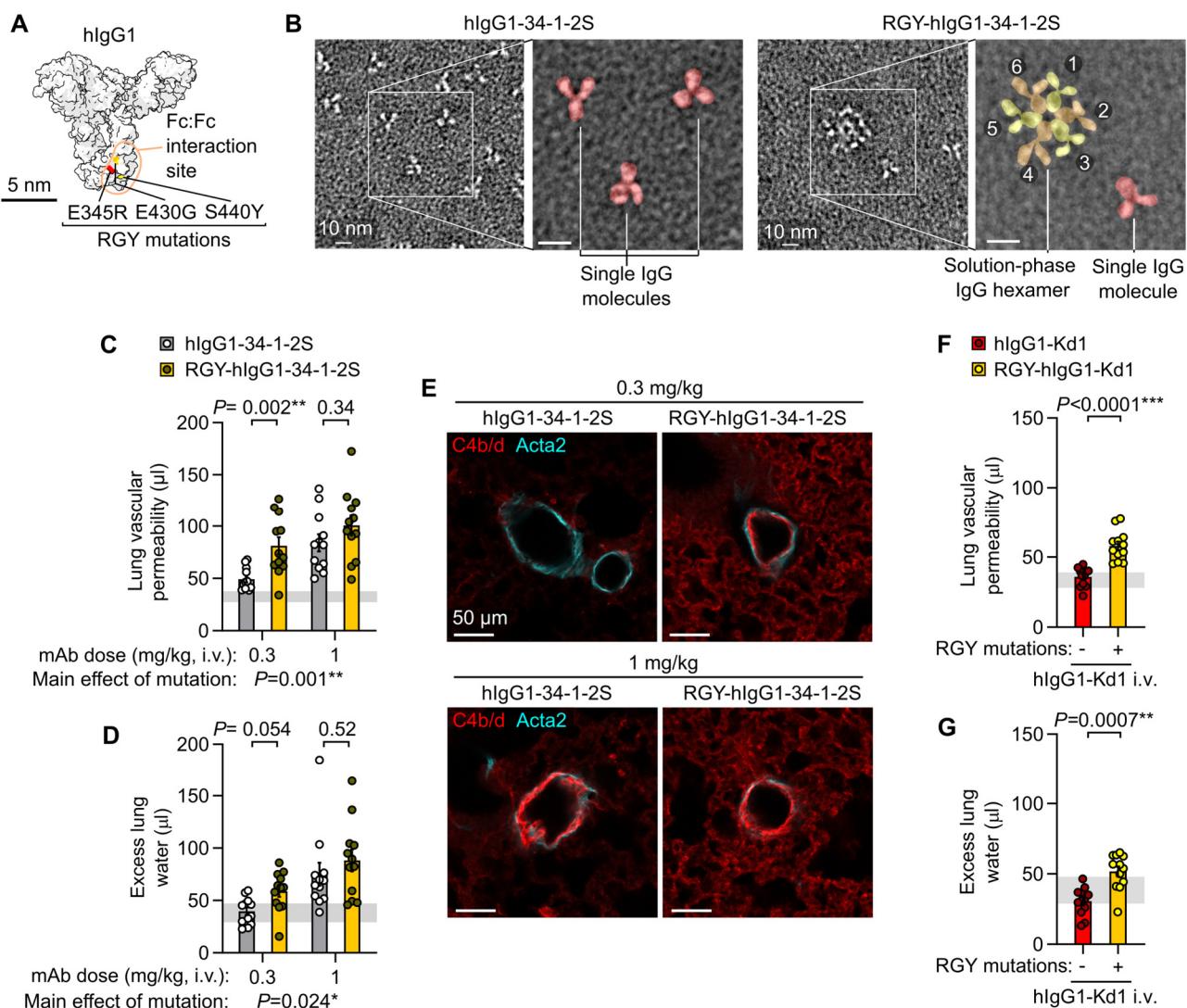
501 **J.** Molecular model showing binding site of SpA-B to Fc domain of human IgG1 (hlgG1), PDB entries 1HZH and 5U4Y.

502 **K.** Lung vascular permeability, **L.** excess lung water measurements and **M.** lung complement C4b/d and Acta2 immunostains from LPS-primed B6.H2d mice after i.v. injection with hlgG1-34-1-2S either mixed with 75 μg SpA-B or vehicle control.

503 Samples for injury measurements were collected at 2 hours after antibody injections and lungs were fixed for immunostaining

504 at 5 minutes after antibody injections. Graphs show means ± standard errors. P-values are from unpaired two-tailed t-tests on

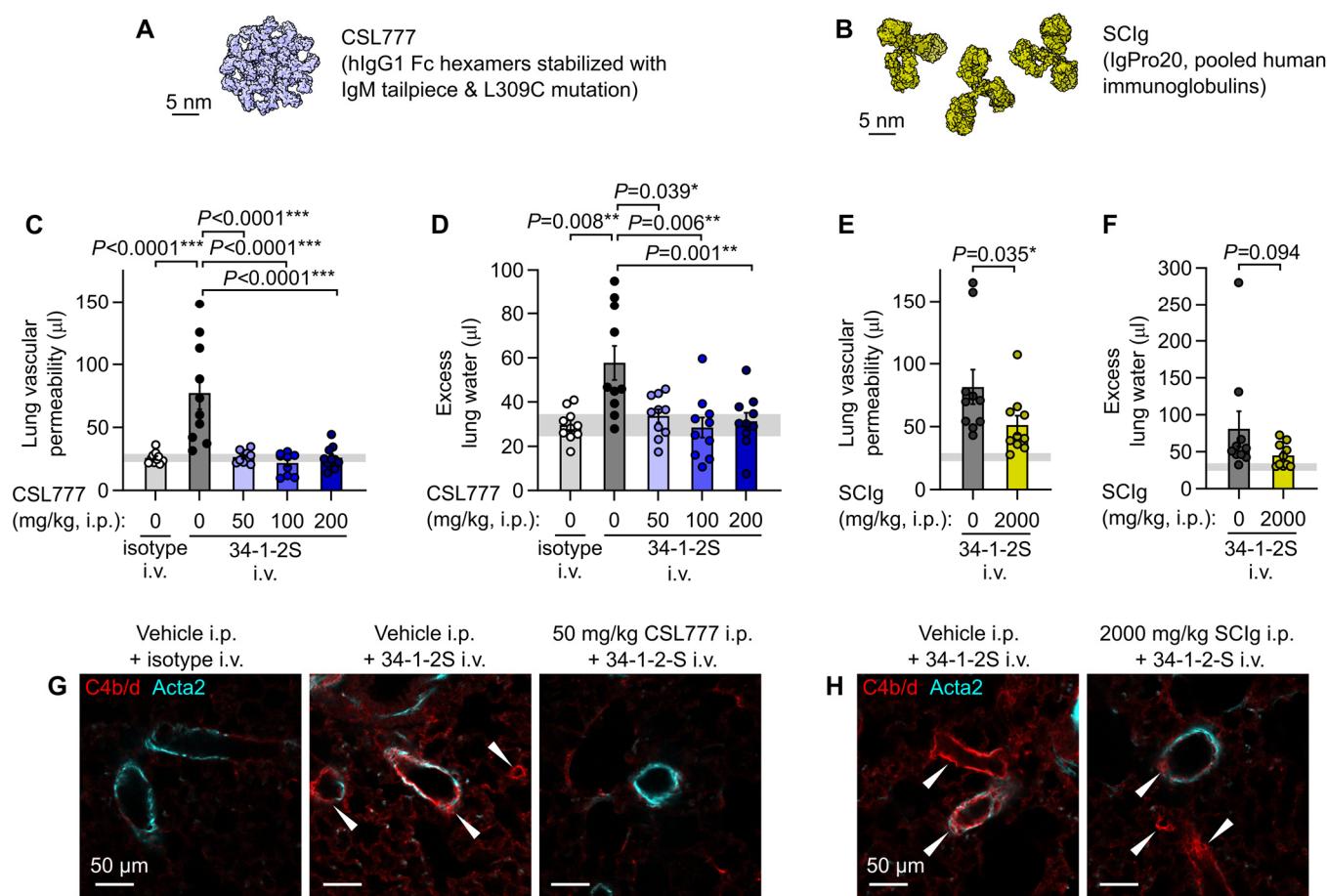
505 log₁₀-transformed data, with group n=8 (**C**, **D**) or n=12 (**G**, **H**, **K**, **L**).



509

510 **Fig. 4. Fc mutations promoting IgG hexamer assembly increase in vivo pathogenicity of alloantibodies.**

511 **A.** Molecular model showing amino acids mutated in RGY-hIgG1 antibodies, based on PDB entry 1HZH.
512 **B.** Negative stain electron micrographs showing single hIgG1-34-1-2S molecules and spontaneous solution-phase hexamers
513 formed by RGY-hIgG1-34-1-2S (colored overlay highlights structures in expanded images).
514 **C.** Lung vascular permeability and **D.** excess lung water measurements from LPS-primed B6.H2^d mice injected with control or
515 RGY-mutated hIgG1-34-1-2S monoclonal antibodies (mAbs) at i.v. doses of either 0.3 or 1 mg/kg.
516 **E.** Immunofluorescence staining showing pulmonary arterioles stained for complement C4b/d (red) and Acta2 (cyan) in lung
517 sections from LPS-primed B6.H2^d mice given indicated treatments, with samples fixed 5 minutes after antibody injections.
518 **F.** Lung vascular permeability and **G.** excess lung water measurements from LPS-primed B6.H2^d mice injected with control or
519 RGY-mutated hIgG1-Kd1 (a novel mAb targeting only the H-2K^d MHC class I antigen) at 1 mg/kg i.v..
520 Graphs show means \pm standard errors with horizontal line representing standard deviations from 'no injury' controls (LPS-
521 primed B6.H2^d mice given hIgG1 isotype control i.v.). Log₁₀-transformed data were analyzed using an ordinary two-way
522 ANOVA with Šídák's multiple comparisons test for effect of Fc mutation within dose level (**C**, **D**) or unpaired two-tailed t-test (**F**,
523 **G**), with group n=12.

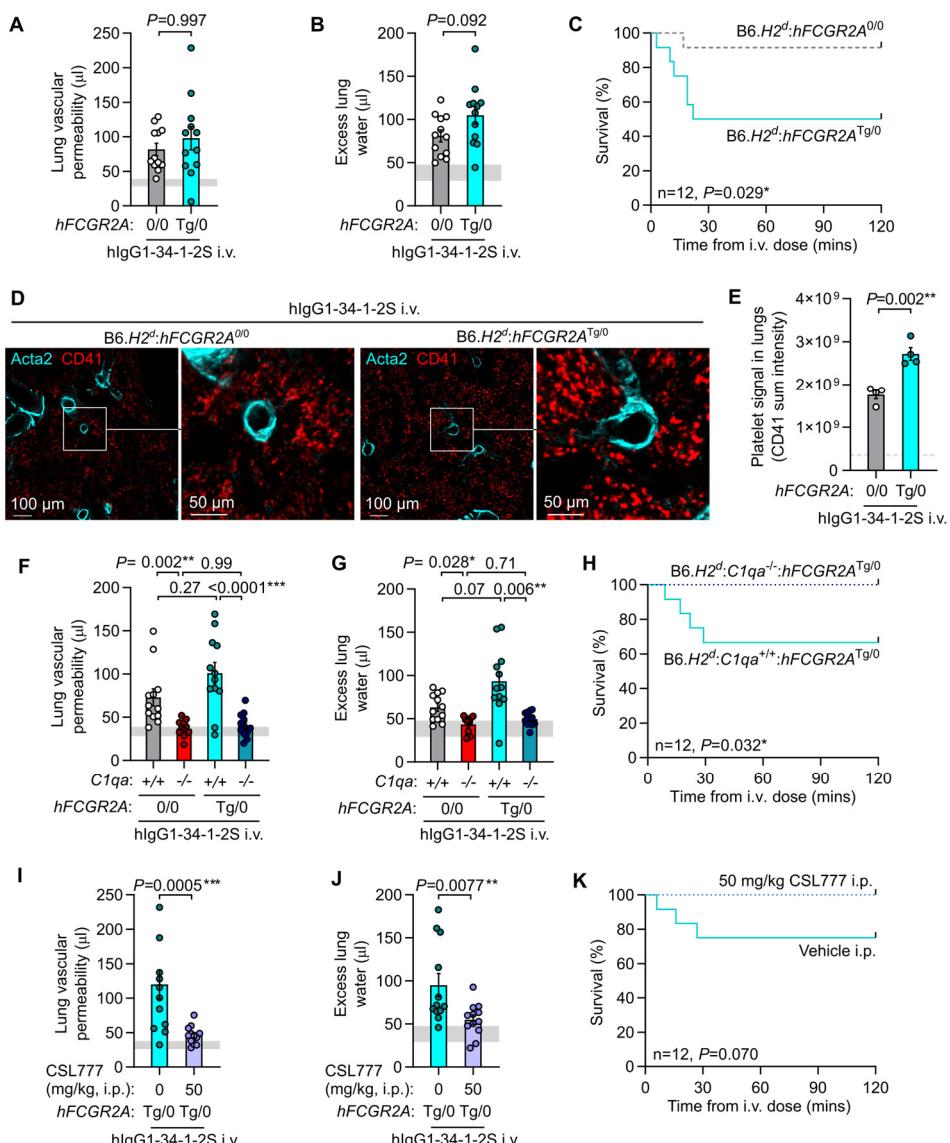


524

525 **Fig. 5. Treatment with recombinant Fc hexamer decoys prevents alloantibody-mediated acute lung injury.**

526 **A.** Molecular representation of CSL777, an investigational recombinant Fc hexamer 'decoy' treatment which inhibits classical
527 complement activation, based on PDB entry 7X13 (50).
528 **B.** Molecular representation of SC Ig, a pooled human immunoglobulin therapeutic with anti-inflammatory properties at high
529 doses, based on PDB entry 1HZH.
530 **C.** Lung vascular permeability and **D.** excess lung water measurements from LPS-primed BALB/c mice given i.p. vehicle or
531 CSL777 at indicated doses 1 hour before i.v. injection with 34-1-2S or mlgG2a isotype control.
532 **E.** Lung vascular permeability and **F.** excess lung water measurements from LPS-primed BALB/c mice given i.p. vehicle or
533 2000 mg/kg SC Ig 1 hour before i.v. injection with 34-1-2S or mlgG2a isotype control.
534 **G.** and **H.** Immunofluorescence showing pulmonary arterioles stained for complement C4b/d (red) and Acta2 (cyan) in lung
535 sections from LPS-primed BALB/c mice given indicated treatments, with samples fixed 5 minutes after antibody injections.
536 White arrowheads point to arterioles with endothelial positivity for C4b/d.
537 Graphs show means \pm standard errors with horizontal line representing standard deviations from 'no injury' controls (from
538 vehicle + isotype control group). Log₁₀-transformed data were analyzed using either (**C**, **D**) an ordinary one-way ANOVA with
539 *P*-values from Dunnett's multiple comparisons test for difference relative to vehicle + 34-1-2S group, or (**E**, **F**) two-tailed
540 unpaired t-test, with group n=10.

541



542

543 **Fig. 6. Acute lung injury is complement-dependent in a model incorporating human FCGR2A-mediated pathology.**
 544 **A.** Lung vascular permeability, **B.** excess lung water and **C.** survival readouts from LPS-primed B6.H2^d:hFCGR2A^{Tg/0} mice and
 545 B6.H2^d littermate controls given i.v. hlgG1-34-1-2S at 1 mg/kg.
 546 **D.** Immunofluorescence imaging of platelet sequestration (CD41, red, with Acta2 in cyan) in lungs from B6.H2^d:hFCGR2A^{Tg/0}
 547 mice and littermates without hFCGR2A fixed at 20 minutes after hlgG1-34-1-2S injections, quantified in **E**.
 548 **F.** Lung vascular permeability, **G.** excess lung water and **H.** survival readouts from LPS-primed B6.H2^d:C1qa^{+/+} and
 549 B6.H2^d:C1qa^{-/-} mice, as well as littermates of each genotype expressing hFCGR2A, given i.v. hlgG1-34-1-2S at 1 mg/kg.
 550 **I.** Lung vascular permeability, **J.** excess lung water and **K.** survival readouts from LPS-primed B6.H2^d:hFCGR2A^{Tg/0} mice given
 551 either vehicle or 50 mg/kg CSL777 i.p. before i.v. hlgG1-34-1-2S at 1 mg/kg.
 552 **A, B, E, F, G, I & J** show means \pm standard errors with horizontal gray lines showing means or standard deviations of values
 553 from 'no injury' controls (B6.H2^d mice given LPS i.p. + hlgG1 isotype control i.v.) and were log₁₀-transformed prior to analysis.
 554 P-values are from: **(A, B, I, J)** unpaired, two tailed t-tests; **(F, G)** two-way ANOVA with Šídák's multiple comparisons test; or
 555 **(C, H, K)** log-rank test, with group n=4 (**E**) or 12 (other graphs).