Role of Repetitive Antigen Patterns for Induction of Antibodies Against Antibodies

By Thomas Fehr,* Martin F. Bachmann,* Etienne Bucher,* Ulrich Kalinke,* Franco E. Di Padova,‡ Alois B. Lang,§ Hans Hengartner,* and Rolf M. Zinkernagel*

From the *Institute of Experimental Immunology, University of Zürich, CH-8091 Zürich, Switzerland; ‡Novartis Pharma AG, S-386.110, CH-4002 Basel, Switzerland; and the §Swiss Serum and Vaccine Institute, Department of Immunology, CH-3001 Bern, Switzerland

Summary
Antibody responses against antibodies, such as rheumatoid factors, are found in several immunopathological diseases and may play a role in disease pathogenesis. Experience shows that they are usually difficult to induce experimentally. Antibodies specific for immunoglobulin constant regions (anti-allotypic) or for variable regions (anti-idiotypic) have been investigated in animal models; the latter have even been postulated to regulate antibody and T cell responses via network-like interactions. Why and how such anti-antibodies are induced during autoimmune diseases, has remained largely unclear. Because repetitively arranged epitopes in a paracrystalline structure of a viral envelope cross-link B cell receptors efficiently to induce a prompt T-independent IgM response, this study used immune complexes containing viruses or bacteria to evaluate the role of antigen pattern for induction of anti-antibody responses. We present evidence that antibodies bound to strictly ordered, but not to irregularly arranged, antigens dramatically enhance induction of anti-antibodies, already after a single immunization and without using adjuvants. The results indicate a novel link between anti-antibody responses and infectious agents, and suggest a similar role for repetitive self-antigens such as DNA or collagen involved in chronic immunopathological diseases.

Abbreviations used in this paper: HRPO, horseradish peroxidase; IC, immune complex; P.a., Pseudomonas aeruginosa; RF, rheumatoid factors S.t., Salmonella typhi; TS, trypic soy; VSV, vesicular stomatitis virus; VSV-G, VSV glycoprotein; VSV-G-huH1, fusion protein of VSV-G with human IgG1 heavy chain constant regions; VSV-IND, VSV serotype Indiana; VSV-NJ, VSV serotype New Jersey.
Materials and Methods

Infectious Agents. VSV serotype Indiana, (VSV-IND, M udd Summers isolate) and VSV serotype New Jersey, (VSV-NJ, Pringle isolate) were originally obtained from Professor D. Kolakowsky (University of Geneva, Switzerland) and grown on BHK cells in minimal essential medium. UV inactivation was performed as described earlier (22) and monitored by a plaque assay on Vero cells. Recombinant VSV-G protein was produced in a baculovirus expression system as described (23); recombinant baculovirus expressing VSV-G was a gift from Dr. D.H.L. Bishop (NERC Institute of Virology, Oxford, UK). Salmonella typhi strain E.83.728 was provided by F. Sadallah (University of Geneva, Switzerland).

Pseudomonas aeruginosa strain Fischer IT-2 was obtained from the Swiss Serum and Vaccine Institute. Both bacteria were grown in tryptic soy (TS) broth at 37°C, quantified on TS agar plates and inactivated as a thin layer in a petri dish by UV irradiation for 10 min (Philips UV lamp, 15 W, distance 8 cm).

Antibodies and IC. Anti-VSV mAb were obtained by fusion of a VSV-immune spleen from BALB/c mice on day 4 after primary (for IgM-secreting hybridomas) or on day 4 after secondary infection (for IgG-secreting hybridomas). The antibodies WN1 222-5 and WN 4 245-2 (both IgG2a) are broadly reactive anti-LPS-core antibodies derived from NZB mice (24). The antibodies 99-T2 (IgG2b) and 63-T2 (IgM) are highly specific anti-LPS-O-chain antibodies against Pseudomonas aeruginosa strain Fisher IT-2 and were generated in BALB/c mice (25). IC were generated by incubation of a mixture of UV-inactivated virus or bacteria with the respective antibodies for 1 h at room temperature. IC formation in the VSV model could be demonstrated indirectly by reduction of anti-VSV neutralizing antibody titers in mice immunized with IC compared with mice immunized with VSV alone.

ELISA for Anti-antibody Detection. We used a sandwich ELISA with the following steps: (a) coating with isotype-specific goat anti-mouse antibody (1 μg/ml; Southern Biotechnologies, Birmingham, AL), (b) blocking with 2% BSA (Fluka, Buchs, Switzerland) in PBS, (c) mAb supernatant (0.2 μg/ml), (d) 20-fold prediluted mouse serum, titrated 1:2 over 11 dilution steps, (e) isotype-specific horseradish peroxidase (HRPO)–labeled goat anti-mouse antibodies (0.5 μg/ml, Southern Biotechnologies), (f) substrate ABTS (2,2′-azino-di-[3-ethylbenzthiazolin-sulfonate (6)], Boehringer Mannheim) and H₂O₂ (Fluka). Plates were coated over night at 4°C, all other incubations were for 60 to 90 min at room temperature. Between incubations, plates were washed three times with PBS containing 0.5 ml TWEEN 20 per liter. OD was measured at 405 nm in an ELISA reader.

All anti-antibody titers are indicated as -log₂ of 20-fold prediluted sera. For Fig. 1, C and D, the dilution step at half maximal OD was determined as shown in Fig. 1 A and then taken as anti-antibody titer. For Fig. 1 E, isotype-specific-anti-ICG antibodies (Southern Biotechnologies) were used for coating and detection at the same concentrations as described above.

VSV Neutralizing Antibodies. VSV-IND neutralizing antibodies (Fig. 2) were determined on the indicated timepoints by a plaque reduction assay on Vero cells as described before (26). Titers are indicated as -log₂ of 40-fold prediluted sera.

VSV-G–huH₁. VSV-G–huH₁ fusion protein was generated following procedures developed by Traunecker and Karl-Jalainen (27) and will be described elsewhere (Bucher, E., U., et al.).
Bodies were induced. This might be explained as follows: (a) a strictly repetitive paracrystalline order of antigen in a viral envelope with a spacing of 5 to 10 nm has been shown to facilitate B cell responses even to self-antigen (28). To evaluate whether antibodies bound to such highly organized antigens might also be presented in an ordered and repetitive fashion and therefore could induce anti-antibodies, BALB/c mice were immunized with IC of autologous antibodies bound to vesicular stomatitis virus (VSV) particles exhibiting paracrystalline strictly repetitive glycoproteins (G) in their envelope. Control groups were immunized with irregularly complexed or monomeric antibodies. While in vitro-generated IC containing 1 μg of a monoclonal IgM antibody M1 against VSV-G and 10^6 PFU of UV-inactivated VSV particles efficiently induced anti-antibodies, IC with the same amount of antibody M1 either irregularly complexed with the recombinant VSV-G protein spontaneously aggregating in tail-to-tail micelles (28a) or cross-linked with a rat anti–mouse Cγ1 antibody did not (Fig. 1 A). Similar results were obtained in A/J and C57BL/6 mice, but in this situation, IgG anti-IgM antibodies could also be induced by the poorly organized IC containing recombinant VSV-G, although at much lower titers than with the highly repetitive complex (Fig. 1 B). When BALB/c mice were boosted once after 14 d, only those treated with virus/antibody complexes exhibited IgM-specific IgG titers of 1:3,000, which did not drop significantly over a period of 80 d (Fig. 1 C). They were dependent on the antibody dose used for generation of IC, as shown for two different monoclonal IgM (Fig. 1 D). In similar experiments using different VSV-specific monoclonal IgG instead of IgM antibodies, no anti-antibodies were induced. This might be explained as follows: (a) the potentially immunogenic variable domains of IgG molecules binding to the virus surface are not easily accessible to B cells, or (b) clearance and processing of IgG- and IgM-containing IC is distinct, because the former, but not the latter, exhibit easily accessible Fc domains that could bind to Fc receptors of macrophages, which would then lead to faster clearance of the IC. Also VSV infection itself did not induce anti-antibodies, probably because the virus only abortively replicates extraneuronally in mice and is rapidly eliminated within 1 d (29), before neutralizing IgM antibodies, which could lead to IC formation, are measured.

Specificity of anti-antibodies induced by VSV-IgM complexes. To determine the specificity of these anti-antibodies, we immunized BALB/c mice (IgMγ) or A/J mice (IgMγ) with IC formed with six different monoclonal VSV neutralizing IgM (M1–M6), which were isolated from BALB/c mice. M3 is specific for the distinct serotype VSV-N (New Jersey (VSV-NJ)), the others for VSV Indiana (VSV-IN D). Analysis of these sera on ELISA plates coated with M1 to M6 revealed that BALB/c mice produced anti-antibodies exclusively specific for the IgM used for immunization, whereas sera of A/J mice recognized any IgM as long as they were of BALB/c origin (Table 1). A monoclonal IgM of A/J origin as well as IgM of normal A/J serum was not recognized by either of the sera (data not shown). Therefore, the anti-antibodies induced in BALB/c mice are idiotype specific, whereas those induced in A/J mice were allotype-specific.

One antibody (M2) failed to induce anti-antibodies (Table 1, Experiment 1); this correlated with its 10–100-fold lower neutralizing capacity and IC formation in vitro compared with the other IgM (data not shown). Both BALB/c antiidiotypic and A/J anti-idiotype antibodies were IgGs and, therefore, T helper cell dependent; this is demonstrated by the failure of CD4+ T cell-depleted or of athymic nude mice to respond with anti-antibodies of IgG isotype (Table 1, Experiment 3). Anti-idiotype antibodies were also found in C57BL/6 mice (IgMγ), and in this case we were even able to detect an IgMα allotype-specific response on day 4 after primary immunization with IC (titers of 1:100), but not after immunization with virus or antibody alone (Fig. 1 E). This response could be prolonged by additional use of LPS (data not shown).

Next, it was tested whether the specificity of induced anti-antibodies as shown in ELISA could also be demonstrated in an independent in vivo read out. For this purpose, A/J and BALB/c were immunized twice with IC containing 10^6 PFU inactivated VSV-N and 5 μg of the antibody M3 to induce anti-antibodies. 10 d later these mice were treated with a specific IgM anti-VSV-IN D antibody (M5), that did not cross-react with VSV-NJ. This enabled us to measure the half-life of M5 by determining the neutralizing titer of the sera against VSV-IN D (Fig. 2). In A/J mice, the half-life of M5 was reduced from 28 to 4 h, whereas in BALB/c mice it was comparable to unimmunized controls; this confirmed the presence of antibodies against the constant region of BALB/c immunoglobulin in A/J, but not in BALB/c mice. Thus, in this model situation anti-idiotype antibodies could be shown to mediate serum IgM antibodies by forming IC in vivo. We were not able to use the same experimental approach to assess modu-
Induction of Anti-antibodies by Repetitive Immune Complexes

To evaluate whether these findings hold true not only for VSV, but also for bacteria exhibiting highly repetitive antigens on their surface, IC formed with gram negative bacteria (exhibiting regularly spaced LPS molecules) and anti-LPS antibodies were tested. $10^8$ CFU of *Salmonella typhi* (**S.t.**) (Fig. 3A) or *Pseudomonas aeruginosa* (P.a.) (Fig. 3B) were UV-inactivated and complexed with a monoclonal IgG2a (origin: NZB) or IgG2b (origin: BALB/c) anti-LPS antibody, respectively, to immunize BALB/c mice. After 14 d, these mice were boosted once, and 6 d later IgG anti-IgG antibodies were found in both situations. The induction of these anti-antibodies was again dependent on the antibody dose used to form IC, as shown for S.t. (Fig. 3A). With the highest dose ($10^8$ CFU) immunization with bacteria alone also induced some anti-antibodies, but at much lower titers than with IC. To test whether these results reflected polyclonal B cell activation by the bacterial LPS, we immunized LPS-responder (C3H/HeN, BALB/c) and LPS-nonresponder (C3H/HeJ) mice with IC or noncomplexed antibodies (Fig. 3D) and tested the BALB/c sera on different antibodies of the same isotype (Fig. 3C). The results indicate: (a) Efficient LPS-mediated stimulation of B cells in the case of P.a., which, however, depended on the presence of IC; the induced anti-antibodies were of rheumatoid factor type because they recognized any autologous IgG2b of BALB/c origin (Fig. 3C). (b) Successful induction of specific anti-allotypic antibodies by IC with S.t., which recognized only heterologous NZB-derived IgGs.
and were independent of B cell activation by LPS (Fig. 3 D). In the case of P.a., 38% (18 of 48) of the mice immunized twice with identical complexes died upon the second immunization from a shock-like syndrome compared with only 4% (2 of 48) of the control groups that were immunized with bacteria, purified LPS, or antibody only; this in vivo finding confirmed the presence of anti-antibodies measured in vitro and, in addition, suggested a pathogenic function of RF-like anti-antibodies.

Immunogenicity of Multivalent Versus Bivalent IC. The notion that antibodies bound to repetitively ordered viral or bacterial antigens induced anti-antibodies was further tested with a sort of inversed IC. Complexes were formed between a VSV-G–specific decavalent IgM (M1) or a bivalent IgG2a antibody (VI49; reference 30) as core of the complex that binds a fusion protein of VSV-G plus constant part of the human IgG1 H chain (huHg1) molecule. These IC display the Fc portions of huIgG1 as repetitive domains and form under optimal conditions decameric (with IgM) or dimeric (with IgG) complexes. In primary and secondary immune responses of BALB/c mice, the fusion protein complexed with the decavalent IgM induced much higher titers of anti-antibodies to huHg1 than bivalent IgG-complexed or the noncomplexed fusion protein alone (Fig. 4). These anti-antibodies were of IgG isotype and T-help dependent, as shown by the negative effect of in vivo CD4+ T cell depletion. Although a very rigid IC structure cannot be assumed in this model, the results show that antibody responses to a foreign constant IgG region, which involves species differences, can be markedly enhanced by multimeric aggregation compared with dimers or monomers. Importantly, they may even suggest that IgM-bound antigen in turn binding IgG may be able to induce anti-IgG antibodies.

In conclusion, the presented data suggest that anti-antibodies may be induced by antibodies bound to highly ordered repetitive antigens, but not by antibodies bound to oligo- or monomeric antigens. Formation of IC was necessary for induction of anti-allootypic, anti-idiotypic and RF-
Our findings indicate a novel link between anti-antibody responses and infectious agents or repetitive self-antigens. The role of highly ordered multimeric antigen patterns for efficient B cell activation has been tested systematically in several model situations involving linear (flagellin [35]; haptenated polymers [36]) or particulate antigens (H antigen B core antigen [37] or VSV [26]). Most infectious agents including viruses, bacteria, and parasites expose highly ordered repetitive antigenic epitopes on their surface. They are able to induce efficient T-independent IgM responses by cross-linking B cell receptors (21, 26, 38, 39). This study now demonstrates that the same rules seem to apply for induction of anti-antibodies. These anti-antibodies were readily induced after only two injections of repetitive IC involving mAbs, but without using adjuvants. In view of the observation that protective anti-Hemophilus influenzae (40) and anti-viral antibodies (30) are of restricted or virtually monoclonal specificity, our results with mAbs against infectious agents can probably be generalized. These findings may explain the secretion of RF during virus infections (influenza [41], rubella, CMV, EBV, HIV) (6) and how the same configuration together with or without LPS may sustain anti-antibody responses in the course of bacterial infections (such as bacterial endocarditis, tuberculosis, or syphilis; for review see 42). Especially chronic infections with persistence of the pathogen during an ongoing antibody response would fulfill the conditions for IC formation in vivo, as it is described for herpesviruses (43, 44), HIV (45, 46), or mycobacteria (47, 48). In addition, the results in BALB/c mice could suggest that, instead of RF, anti-idiotypic antibodies may enhance immunopathology found in reactive forms of arthritis, which are apparently RF negative. Also, repetitive self antigens (e.g., collagen II [49] in rheumatoid lesions, DNA [50] in SLE) exposed after primary or secondary tissue injury may induce anti-antibodies by formation of highly repetitive immune complexes. This may explain why RF secretion is such a typical manifestation accompanying these diseases (15).

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Address correspondence to R. M. Zinkernagel, Institute of Experimental Immunology, University of Zürich, Schmelzbergstrasse 12, CH-8091 Zürich, Switzerland. Dr. Bachmann's present address is Ontario Cancer Institute, Department of Medical Biophysics and Immunology, University of Toronto, Canada M5G 2M9.

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References


