Efficient Neutralization of Anthrax Toxin by Chimpanzee Monoclonal Antibodies against Protective Antigen

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Four single-chain variable fragments (scFvs) against protective antigen (PA) and 2 scFvs against lethal factor (LF) of anthrax were isolated from a phage display library generated from immunized chimpanzees. Only 2 scFvs recognizing PA (W1 and W2) neutralized the cytotoxicity of lethal toxin in a macrophage lysis assay. Full-length immunoglobulin G (IgG) of W1 and W2 efficiently protected rats from anthrax toxin challenge. The epitope recognized by W1 and W2 was conformational and was formed by C-terminal amino acids 614–735 of PA. W1 and W2 each bound to PA with an equilibrium dissociation constant of $10^{-11}$ mol/L, which is an affinity that is 20–100-fold higher than that for the interaction of the receptor and PA. W1 and W2 inhibited the binding of PA to the receptor, suggesting that this was the mechanism of protection. These data suggest that W1 and W2 chimpanzee monoclonal antibodies may serve as PA entry inhibitors for use in the emergency prophylaxis against and treatment of anthrax.

Anthrax has emerged as a serious bioterrorist threat. Inhalational anthrax is usually fatal if treatment is delayed [1]. The lethality of anthrax is primarily the result of the effects of anthrax toxin, which has 3 components: a receptor-binding protein known as “protective antigen” (PA) and 2 catalytic proteins known as “lethal factor” (LF) and “edema factor” (EF). Entry of the toxins into the cell is initiated by rapid binding of the 83-kDa PA to the cellular receptor, whereupon a furinlike protease cleaves the bound PA into an N-ter-

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minal 20-kDa protein (PA20) and a C-terminal 63-kDa protein (PA63) [2]. There is spontaneous oligomerization of PA63 into an antigenically distinct heptameric ring that can no longer be displaced from the cellular receptor. The heptamer binds up to 3 molecules of LF or EF. The resulting complexes enter the cell by endocytosis, and a conformational change induced by low pH results in the release of bound LF and EF into the cytosol. Several steps in this process could be targets for antibodies; for example, antibodies to PA might block receptor binding, oligomerization, or binding of LF and EF, and antibodies to LF and EF might prevent their binding to PA.

Passive immunization with polyclonal antibodies protects laboratory animals from the effects of anthrax toxins [3, 4]. Passive immunization of humans with anthrax neutralizing antibodies may provide an effective treatment when vaccination against anthrax is not practical or antibiotic treatment is not effective. Antibody therapy given in conjunction with antibiotics would also be useful when the accumulated level of toxin would be detrimental, even if further bacterial growth
Figure 1. Alignment of the deduced amino acid sequences of the variable domains of the heavy chain (VH) and \( \kappa \) chain (VK). Substitutions relative to W1 are expressed as single letters denoting amino acids. Dashes denote identical residues, and asterisks denote the absence of corresponding residues relative to the longest sequence. Complementarity-determining regions (CDR1–3) and framework regions (FR1–4) are shown above the sequence alignments.

was inhibited. Recently, several recombinant monoclonal antibodies (MAbs) against PA were shown to protect animals from challenge with anthrax toxin [5–7]. Because chimpanzee immunoglobulins are virtually identical to human immunoglobulins [8–10], high-affinity chimpanzee antibodies that neutralize anthrax toxins should have therapeutic value comparable to that of human antibodies. Here, we report the identification and characterization of potent neutralizing chimpanzee MAbs against PA.

**MATERIALS AND METHODS**

**Reagents.** Recombinant PA, PA63, LF, and EF were obtained from List Biologicals or were made in our laboratory. Enzymes used in molecular cloning were purchased from New England BioLabs, and oligonucleotides were synthesized by Invitrogen. Anti-His horseradish peroxidase (HRP) conjugate, anti–human Fab HRP conjugate, and anti–human Fc agarose were purchased from Sigma. Nickel-agarose beads were obtained from Invitrogen.

**Animals.** Chimpanzees 1603 and 1609 were immunized 3 times, at 2-week intervals, with 50 \( \mu g \) each of recombinant PA, LF, and EF with alum adjuvant. Eight weeks after the last immunization, bone marrow was aspirated from the iliac crest of these animals. Fischer 344 rats were purchased from Taconic Farms. All experiments involving animals were performed under protocols approved by the Animal Care and Use Committee of the National Institute of Allergy and Infectious Diseases, National Institutes of Health.

**Library construction and selection.** Lymphocytes from the bone marrow aspirate were isolated on a Ficoll gradient. mRNA was extracted from lymphocytes by use of an mRNA purification kit (Amersham Biosciences). cDNA was synthesized using a first-strand cDNA synthesis kit (Amersham Biosciences). The heavy-chain (VH) and \( \kappa \) light-chain (VK) genes were amplified by polymerase chain reaction (PCR) for 30 cycles (for 1 min at 95°C, for 1 min at 58°C, and for 1 min at 72°C), by use of a mixture of primers for human V genes described elsewhere [11]. The single-chain variable fragment (scFv) was assembled from VH and VK via splicing by overlap extension PCR. Gel-purified scFv DNA was digested with SfiI and then was cloned into a pAK100 vector that also had been cut with SfiI [12].

The recombinant plasmids were transformed into *Escherichia coli* XL1-blue (Stratagene) by electroporation, resulting in a library of 5 \( \times 10^7 \) individual clones. The phagemid production, panning, and screening were essentially the same as those reported by Krebber et al. [12] in 1997. In brief, the phagemids were rescued by superinfection with helper phage VCS M13 (Stratagene) and then were subjected to panning on PA, LF, and EF proteins coated on ELISA wells. Nonspecifically adsorbed phages were removed by extensive washing. Specifically
bound phages were eluted with 100 mmol/L triethylamine, neutralized in pH, and amplified. After 3 rounds of panning, randomly chosen single-panhe scFv clones were screened for specific binding by phage ELISA [13]. Clones that differentially bound to specific antigens with values of >1.0 for absorbance measured at 450 nm (A450) were considered to have a positive result, whereas values of <0.2 were considered to denote a negative result. For clones that bound specifically to PA, LF, or EF, the variable regions of VH and light (VL)–chain genes were aligned. 

The VH and Vλ, CH1, and Jλ segments were considered to have a positive result. For clones that bound specifically to PA, LF, or EF, the variable regions of VH and light (VL)–chain genes were aligned. Their corresponding amino acid sequences were sequenced, and they were cloned into pComb3H at the matching restriction sites [14].

Conversion of scFv to Fab and to full-length IgG. To convert scFv to Fab, the following 8 primers were designed:

1. Fab H-5′: ACCGTCTCCTCAGCCTCCACCAAGGGCCCATCGGT
2. Fab H-3′: AATGAGATCTCGGCCGCTTAAATTAA
3. Fab L-5′: GTGGAAATCAAACGAACTGTGGCTGCACTCTGTG
4. Fab L-3′: AGGTATTTCATTTTAAATTCCTCCT
5. PA H-5′: AGGTTGAGGCTAGGAGCGAGCTGGACAGGAGGCCATCGGT
6. PA H-3′: CTCCGATCCGAGGTGAGAGGAGGAGCGAGCTGGACAGGAGGCCATCGGT
7. PA L-5′: TGGAGGATCCGAGGCTAGGAGCGAGCTGGACAGGAGGCCATCGGT
8. PA L-3′: AGCCACAGTCTTGTTGATTCCACCTTGCTGTTCCAGG

First-strand cDNA of constant chain 1 of heavy chain (CH1) and constant region of k light chain (Ck) gene fragments was synthesized from total RNA obtained from human spleen cells (BD Biosciences) and was amplified by PCR performed using Fab H-5′/Fab H-3′ and Fab L-5′/Fab L-3′ primers, respectively. The VH and Vx gene fragments were amplified by PCR performed using anti-PA scFv DNA as a template and PA H-5′/PA H-3′ and PA L-5′/PA L-3′ as primers. The Fd (VH-CH1) and k-chain (Vx-Ck) segments were produced through splicing by overlap extension PCR of VH CH1 and of Vx and Ck, with use of PA H-5′/Fab H-3′ and PA L-5′/Fab L-3′ as primers. The Fd region was digested with Xhol and NotI, and the k-chain region was digested with XbaI and SacI. The digested DNA fragments were cloned into pComb3H at the matching restriction sites [14].

Fab was converted to full-length IgG by digestion of Fd with Xhol and Apal and by cloning into a pCDHGC68b vector [15] that contains human heavy-chain constant region to yield plasmid pH11003. The k-chain was digested with XbaI and SacI and was cloned into a pCNHL vector [15] to yield plasmid pH11034.

Expression and purification of scFv, Fab, and IgG. Both scFv and Fab were expressed in E. coli. In brief, bacteria were cultured at 30°C in 2X YT (yeast extract–tryptone–rich) medium containing 2% glucose and appropriate antibiotics, until the OD600 was 0.5–1. The culture was diluted 5-fold in 2X YT without glucose and containing 0.2 mmol/L isopropyl-β-D-thiogalactopyranoside, and it then was incubated for 20 h at 27°C. The expressed proteins were purified by chromatography performed on nickel-charged affinity resins (Invitrogen).

The full-length IgG plasmids pH11003 and pH11034, which contained the VH and VL of anti-anthrax toxin components, were cotransfected into 293T cells for transient expression. The IgG was purified by affinity chromatography performed on human Fc agarose (Sigma).

The purity of each antibody was determined by SDS-PAGE (Novex; Invitrogen), and the protein concentration was determined by bicinchoninic assay (BCA; Pierce Biotechnology) and ELISA performed with a purified human IgG (Jackson ImmunoResearch) as a reference standard.

ELISA. PA, LF, and unrelated proteins (bovine serum albumin [BSA], thyroglobulin, lysozyme, and phosphorylase b) were coated onto a 96-well plate by placing 100 μL of protein in carbonate buffer (pH 9.5) at a concentration of 5 μg/mL in each well and incubating the plate at room temperature overnight. Serial dilutions of soluble scFv, Fab, IgG, or phage were added to the wells, and the plates were incubated for 2 h at room temperature. The plates were washed, and the secondary antibody conjugate (anti-His-HRP, anti-human Fab-HRP, or anti-M13-HRP) was added and incubated for 1 h at room

Table 1. Human immunoglobulin germline genes most closely related to the chimpanzee heavy chain (VH) and k light chain (Vx) of various anti-anthrax protective antigen (PA) and lethal factor (LF) antibodies.

| Monoclonal antibody | VH family | VH segment | D segment | JH segment | Vx family | Vx segment | Jx segment |
|---------------------|------------|------------|-----------|------------|-----------|------------|------------|
| W1                  | VH3        | DP-42      | ND        | J6c        | Vκ II     | DPK-15     | Jκ3        |
| W2                  | VH3        | DP-42      | ND        | J6c        | Vκ II     | DPK-15     | Jκ1        |
| W5                  | VH4        | DP-78      | D2-15     | J5b        | Vκ II     | A2b        | Jκ4        |
| A63-6               | VH3        | DP-47      | D1-14     | J6c        | Vκ I      | HK 102     | Jκ5        |
| F3-6                | VH3        | DP-49      | ND        | J4b        | Vκ I      | HK 102     | Jκ5        |
| F5-1                | VH5        | DP-73      | ND        | J6c        | Vκ I      | HK 102     | Jκ1        |

NOTE. The closest human VH and Vx germline genes were identified by V-BASE database similarity searches [25]. ND, not determined because of a lack of identifiable homologue.
Figure 2. ELISA titration of anti–protective antigen (PA) (A) and anti–lethal factor (LF) (B) single-chain variable fragments (scFvs). Recombinant PA or LF or unrelated proteins, bovine serum albumin (BSA), thyroglobulin, lysozyme, and phosphorylase b were used to coat the wells of an ELISA plate. Wells then were incubated with various dilutions of scFvs. Bound scFv was detected by the addition of peroxidase-conjugated anti-His antibody, followed by tetramethylbenzidine substrate. The anti-PA and anti-LF scFvs did not bind to the unrelated proteins; only BSA is shown as an example. OD, optical density measured at 450 nm.

Figure 3. In vitro neutralization assay. Anti–protective antigen (PA) IgG was mixed with anthrax toxin and was incubated for 1 h at 37°C. The mixture was added to RAW264.7 cells in a 96-well plate and was incubated for 4 h at 37°C. After washing, the cells were stained with MTT dye, and then lysis was performed in a solution containing 0.5% SDS in 90% isopropanol and 0.05 mol/L HCl. The plate was read at an optical density measured at 570 nm, and the cell survival was calculated relative to untreated cells. Results were plotted and analyzed using Prism software (version 4; Graphpad Software). EC50, effective concentration for 50% neutralization.
essentially as described elsewhere [21]. In brief, a mixture of a 
<sup>35</sup>S-labeled peptide and anti-PA W2 was incubated at 4°C

overnight, and immune complexes were collected with protein G–
coupled agarose beads (Amersham Biosciences). The precipitated

complex was washed and subjected to SDS-PAGE. The PA pep

tide was detected by exposing the dried gel to an x-ray film.

**PA binding assays.** RAW264.7 cells [22] were grown, in 6-

well plates, to 90% confluence. Ten microliters of PA (10 μg/ 

mL) was mixed with neutralizing antibody 14B7 [23] or with

W2 at molar ratios of PA to MAb of 1:1 or 1:10 in a final

volume of 200 μL, and it then was incubated for 5 min or 15 

min. PBS was mixed with PA in control samples. The mixture

was added to the cells and was incubated for 20 min at 37°C.

The medium was removed, and cells were washed 5 times with

ice-cold PBS, followed by lysis in radioimmunoprecipitation

assay buffer (1% Nonidet, 0.5% sodium deoxycholate, and

0.1% SDS in PBS) plus EDTA-free complete protease inhibitor

cocktail (Roche Applied Science). Equal amounts of protein

(as measured with BCA; Pierce Biotechnology) were loaded

onto SDS-PAGE gels. Western blot analysis was performed us

ing anti-PA polyclonal antibody 5308 (dilution, 1:3500), which

was developed in our laboratory (Bacterial Toxins and Therap

etics Section, National Institute of Allergy and Infectious

Diseases), and HRP-conjugated goat anti–rabbit IgG secondary

antibody (dilution, 1:2000) (Santa Cruz Biotech).

**In vitro neutralization assay.** An established RAW264.7

cell–based assay was used to determine the in vitro neutraliz

ation activity of antibodies [22, 24]. Results were plotted, and

the effective concentration for 50% neutralization (EC<sub>50</sub>) was

calculated using Prism software (Graphpad Software).

![Image](image-url)

**Figure 4.** Competitive ELISA. Recombinant protective antigen (PA) was coated onto the wells of an ELISA plate. Wells then were incubated with anti-PA W2 Fab at the concentrations indicated. After incubation for 1 h at room temperature, anti-PA W2 Fab was removed from the wells, and mouse anti-PA monoclonal antibodies (MAbs) 14B7 and 2D3 [16, 23] were added to the wells. Bound MAbs were detected by the addition of peroxidase-conjugated anti–mouse antibody, followed by tetrathymethylbenzidine substrate. The binding to PA was calculated by dividing the optical density value in the absence of W2 by the optical density value in the presence of W2. ■, 14B7; ▲, 2D3.

| Table 2. Affinity of anti-antibody to anthrax protective antigen (PA) W1 and W2 antibodies, compared with other human anti-PA monoclonal antibodies (MAbs). |
|--------------------------------------------------|
| Antibody  | K<sub>d</sub> (mol/L) | k<sub>d</sub> (s<sup>-1</sup>) | k<sub>a</sub> (s<sup>-1</sup>) | K<sub>v</sub> (mol/L) | Reference |
|-----------|----------------------|----------------------|----------------------|----------------------|-----------|
| W1        | 2.90 × 10<sup>-8</sup> | 1.15 × 10<sup>-4</sup> | 3.97 × 10<sup>-11</sup> | Present study        |
| W2        | 2.77 × 10<sup>-8</sup> | 1.52 × 10<sup>-4</sup> | 5.49 × 10<sup>-11</sup> | Present study        |
| A/VP-21D9 | 1.8 × 10<sup>-8</sup> | 1.48 × 10<sup>-4</sup> | 8.21 × 10<sup>-11</sup> | [7]                  |
| A/VP-1C6  | 1.85 × 10<sup>-8</sup> | 1.31 × 10<sup>-4</sup> | 7.11 × 10<sup>-10</sup> | [7]                  |
| A/VP-4H7  | 1.74 × 10<sup>-8</sup> | 2.45 × 10<sup>-4</sup> | 1.41 × 10<sup>-10</sup> | [7]                  |
| A/VP-22G12 | 1.01 × 10<sup>-8</sup> | 5.17 × 10<sup>-4</sup> | 5.12 × 10<sup>-10</sup> | [7]                  |
| 83K7C     | 1.16 × 10<sup>-8</sup> | 4.26 × 10<sup>-4</sup> | 3.67 × 10<sup>-9</sup>  | [6]                  |
| 63L1D     | 1.50 × 10<sup>-8</sup> | 1.90 × 10<sup>-4</sup> | 1.3 × 10<sup>-10</sup> | [6]                  |
| scFv1     | NA                   | NA                   | 1.9 × 10<sup>-7</sup>  | [26]                 |
| scFv4     | NA                   | NA                   | 3.1 × 10<sup>-7</sup>  | [26]                 |
| scFv12    | NA                   | NA                   | 1.1 × 10<sup>-6</sup>  | [26]                 |
| scFv24    | NA                   | NA                   | 4.3 × 10<sup>-7</sup>  | [26]                 |
| ETI-204<sup>a</sup> | 4.57 × 10<sup>-8</sup> | 1.50 × 10<sup>-4</sup> | 3.3 × 10<sup>-10</sup> | [27]                 |

**NOTE.** K<sub>v</sub>, equilibrium dissociation constant; k<sub>d</sub>, dissociation rate; k<sub>a</sub>, association rate; NA, not available; scFv, single-chain variable fragment.

<sup>a</sup> ETI-204 is an affinity-improved, humanized MAb derived from murine MAb 14B7.
intravenously with PBS or antibody at 5 min, 4 h, or 1 week before an intravenous injection with PA + LF (7.5 μg each).

RESULTS

Isolation of anti-PA and anti-LF clones. Recombinant proteins were used as antigens to select antibodies from an scFv library derived from immunized chimpanzees. After 3 rounds of selection with PA, LF, or EF, a total of 192 phagemid clones were screened for specific binding by use of ELISA. We did not detect clones that bound specifically to EF protein. Ninety percent of the clones recognized PA or LF proteins but not BSA control protein. Four unique anti-PA and 2 unique anti-LF scFv phagemid clones were identified (W1, W2, A63-6, W5, F3-6, and F5-1) by sequence analysis (figure 1). Two clones, W1 and W2, had very similar sequences in the VH and Vk regions (figure 1), with only a 9-aa difference (3 of the amino acids are located within the primer region). The amino acid sequences of the other 4 clones differed greatly. As expected, the greatest divergences in sequence and length were noted in the CDR regions. Because there is extensive homology between chimpanzee and human immunoglobulins [8], similarity searches of all known human immunoglobulin genes in the V-DBASE database were conducted [25]. The VH3 family of VH germ line genes. The 6 scFvs were expressed in E. coli, purified by affinity chromatography, and tested for binding activity and specificity by ELISA. Clones W1, W2, A63-6, and W5 bound strongly to PA antigen, and clones F3-6 and F5-1 bound strongly to LF antigen (figure 2); none of the clones bound to BSA, thiglobulin, phosphorylase b, or lysozyme (data not shown).

In vitro neutralizing activity and affinity of W1 and W2. The effect of anthrax toxin was strongly inhibited by the W1 and W2 scFvs, but not by A63-6, F3-6, F5-1, and W5 scFvs (data not shown). The latter scFvs did not undergo further study.

scFv fragments have limited use in passive immunotherapy, because these monovalent fragments are rapidly cleared from the blood. In most cases, bivalent full-length immunoglobulin is more effective than is the corresponding scFv, because of avidity effects, effector functions, and prolonged half-life in the blood. Therefore, W1 and W2 scFvs were converted to bivalent whole IgG1s and were compared for neutralization activity in the RAW264.7 cell-based in vitro assay. Well-characterized mouse anti-PA 14B7 was used as a comparison control [23].

Neutralization by complete IgGs was 5–20-fold better than neutralization by scFvs (data not shown) and was 5-fold higher (for W2) and 15-fold higher (for W1) than that by mouse anti-PA 14B7 (figure 3). The equilibrium dissociation constant (Kd) for W1 and W2 IgG1, respectively, was determined by Biacore analysis. W1 and W2 antibody displayed very high affinity, with a Kd of 4 × 10⁻¹¹ and 5 × 10⁻¹¹ mol/L, respectively, compared with a Kd of 4 × 10⁻⁸ mol/L for 14B7 (in the present study) (table 2). These affinities compared favorably with those reported for human anti-PA MAbs (table 2), and they were the highest recorded affinities against anthrax PA [6, 7, 26].

Characterization of the neutralization epitope and mechanism of neutralization. Competitive ELISA results indicated that W1 and W2 may recognize the same epitope, because they competed with each other in binding to PA (data not shown), and because the complementarity-determining regions of their VH chains are identical. Therefore, only W2 was used to map the neutralization epitope. In ELISA, competition was also observed between 14B7 and W2 but not between 2D3 and W2 (figure 4). 14B7 binds to the site responsible for binding to the cellular receptor, whereas 2D3 does not compete with 14B7 and binds to a distinct binding site [16]. The binding site for W2 was mapped by radioimmunoprecipitation assay. The smallest peptide that reacted with W2 antibody contained 121 aa, corresponding to aa 614–735 at the C-terminus of PA (figure 5). This result indicated that W2 bound to a conformational epitope that encompassed almost the entire domain 4 of PA [28].
Table 3. Neutralization of protective antigen (PA) of anthrax toxin by W2 in rats.

| Antibody | Molar ratio of MAb to PA | No. of rats that survived/ no. of rats that received injections |
|----------|--------------------------|--------------------------------------------------------------|
| None     | 0/5                      |                                                              |
| 14B7     | 1:4                      | 1/3                                                          |
| W2       | 1:4                      | 3/3                                                          |
| 14B7     | 1:3                      | 3/3                                                          |
| W2       | 1:3                      | 3/3                                                          |

**NOTE.** Antibody and lethal toxin (protective antigen + lethal factor) were mixed at the molar ratios indicated and were injected intravenously. MAb, monoclonal antibody.

- The amount of PA was kept constant (at 7.5 μg).
- Lethal toxin only was administered (amount injected, 7.5 μg).
- Time to death, 100–134 min.
- Time to death, 128 and 168 min.

Because domain 4 of PA is responsible for cellular receptor binding [29, 30], the results suggested that W2 neutralizes the toxin by blocking binding of PA to the cellular receptor. This conclusion was confirmed in a binding assay, by use of Western blot analysis, which showed that W2 prevented the binding of PA to RAW264.7 cells (figure 6). The affinity ($K_d$) of binding of PA to the receptor is $1 \times 10^{-9}$ mol/L to $5 \times 10^{-9}$ mol/L [31]. Therefore, the affinity of W2 antibody binding to PA is 20–100-fold higher than the affinity of PA for the cellular receptor.

**In vivo animal protection.** Because affinity of anti-PA is strongly correlated with its neutralization activity, it is reasonable to assume that anti-PA W1 and W2 are potent neutralizing antibodies. As has been reported for other anti-PA MAbs [6, 7], one remarkable feature of our antibodies is the very slow dissociation rate, which may provide a significant physiological advantage for toxin neutralization in vivo.

To evaluate the neutralization of PA by W1 and W2 in vivo, we measured protection against anthrax toxin challenge in the Fischer 344 rat model in 2 ways. First, MAb and toxin were mixed at different molar ratios, and the mixtures were injected into 3 rats each, which were then observed for morbidity and death. Injection of 7.5 μg of LT (7.5 μg of PA + 7.5 μg of LF) alone killed all 3 rats within 100–134 min (table 3). W2 antibody conferred protection at very low concentrations. Addition of W2 antibody at a molar ratio of 1:4 (Ab:PA), the lowest concentration of antibody tested, completely protected the rats from anthrax toxin challenge. In comparison, 14B7 mouse antibody protected only 1 of the 3 rats at this ratio, probably reflecting the difference in affinity between W2 and 14B7.

Second, in a more stringent test, the MAb was injected 5 min, 4 h, and 1 week before injection of LT (7.5 μg of PA + 7.5 μg of LF), to investigate the duration of antibody protection. Single administration of W1 and W2 antibodies at a 2:1 molar ratio (Ab:PA) protected 6 of 6 rats challenged with toxin 5 min, 4 h, or 1 week later (table 4). A lower dose of antibodies (0.5:1) still protected all the rats ($n = 6$) when they were challenged 5 min or 4 h later but not when they were challenged 1 week later (table 4).

**DISCUSSION**

We identified 2 chimpanzee MAbs, W1 and W2, which bind to PA with high affinity. These antibodies neutralized cytotoxicity of anthrax toxin in the picomolar range in vitro, and they efficiently protected animals from anthrax toxin challenge in vivo, most likely by blocking binding to the cell receptor. Our 2 neutralizing MAbs have the highest affinity of any human antibodies for PA reported thus far. Antibody affinity has been shown to correlate well with efficacy [5, 32–36].

The neutralization epitope recognized by W1 and W2 MAbs was mapped to a region of PA comprising aa 614–735. So far, 3 neutralization epitopes in PA have been proposed: the site for binding to the cellular receptor, the site for binding to LF [16, 37], and the site for heptamer formation [37]. However, the locations of these putative epitopes have not been precisely defined. For example, the epitope recognized by murine MAB 14B7 was suggested to be between Asp-671 and Ile-721, on the basis of the differential binding of 14B7 to different PA fragments generated through C-terminal deletions [16]. Because N-terminally deleted PA fragments were not tested, the epi-

Table 4. Duration of neutralization of protective antigen (PA) of anthrax toxin by W1 and W2 in rats.

| Molar ratio of MAb to PA | 5 min $^b$ | 4 h $^b$ | 1 week $^b$ |
|-------------------------|------------|----------|-------------|
| W1                      | 3/3        | 3/3      | 3/3         |
| W2                      | 3/3        | 3/3      | 3/3         |
| Combined                | 6/6        | 6/6      | 6/6         |

**NOTE.** LF, lethal factor; MAb, monoclonal antibody.

- Injection with 7.5 μg of PA plus 7.5 μg of LF.
- The interval between injection with MAb and injection with PA plus LF.
Anthrax, whether resulting from natural or bioterrorist-associated exposure, is a constant threat to human health. Although production of an efficient anthrax vaccine is an ultimate goal, the benefits of vaccination can be expected only if a large proportion of the population at risk is immunized. The low incidence of anthrax suggests that large-scale vaccination may not be the most efficient means of controlling this disease. In contrast, passive administration of neutralizing human or chimpanzee MAb to a subject at risk for anthrax or exposed to anthrax could provide immediate efficacy for emergency prophylaxis against or treatment of anthrax. This finding is supported by a recent study [27], which indicated that passive immunization with affinity-improved, humanized murine MAb 14B7 against PA (ETI-204) could efficiently protect rabbits before or after challenge with aerosolized Bacillus anthracis spores. However, humanized murine MAbs may retain some antigenic components of the original murine sequences and may elicit antibodies to the MAb in humans. Human- and chimpanzee-derived MAbs would not be expected to have this problem, because the sequences of chimpanzee immunoglobulin genes derived MAbs would not be expected to have this problem, components of the original murine sequences and may elicit antibodies to the MAb in humans. Human- and chimpanzee-derived MAbs would not be expected to have this problem, because the sequences of chimpanzee immunoglobulin genes are virtually identical to those of humans [8–10, 40]. Furthermore, one would expect that the anti-PA chimpanzee MAbs W1 and W2 will provide better protection than will ETI-204, because these MAbs had higher affinity ($K_d$) and lower EC$_{50}$ in in vitro neutralization experiments than did ETI-204.

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