Inhibition of astrocyte proliferation and binding to brain tissue of anticardiolipin antibodies purified from lupus serum

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Abstract
Polyclonal antihistone antibodies purified from pooled serum samples of patients with systemic lupus erythematosus were shown to have inhibitory effects on cultured normal rat brain astrocytes (RBA-1 cells). Antihistone antibodies at concentrations from 50 to 200 μg/ml inhibited the [3H]thymidine incorporation of RBA-1 cells in a dose dependent manner. A kinetic study showed that antihistone antibodies (100 μg/ml) maximally inhibit the proliferation of RBA-1 cells (20-6 (5-1)% of the control value) after incubation for one day. In contrast, human γ globulin (100 μg/ml) had no effect on these cells. In the presence of antihistone antibodies (100 μg/ml), the RBA-1 cells attached to the bottom of wells became spherical and the expression of glial fibrillary acidic protein in the cytoplasm was slightly reduced. Using 3,3′-dihexylxocarbocyanine iodide as an indicator, antihistone antibodies depolarized the membrane potential of RBA-1 cells after one day of culture. In addition, the percentage binding of RBA-1 cells with antihistone antibodies was greater than with γ globulin as determined by flow cytometric analysis. Immunofluorescence staining of brain tissue from BALB/c mice with antihistone antibodies was noted in the corpus callosum, the cellular zone near the corpus callosum, and cells scattered in brain tissue. These results suggest that antihistone antibodies have an inhibitory effect on brain cells and elicit thrombus formation in brain vessels, which plays a part in neuropyschiatric lupus.

Neuropsychiatric manifestations are common in patients with systemic lupus erythematosus (SLE). More than 25% of these patients develop central nervous system disease in the first two years after the diagnosis of SLE. Microscopic studies of the brains of patients with SLE who died have emphasised the presence of widely scattered microinfiants and a non-inflamatory vasculopathy. True vasculitis with inflammatory cells invading vessel walls and fibrinoid necrosis are rarely seen in the nervous system of patients with SLE, however. It is believed that autoantibodies reactive with the cell membrane in the circulation or cerebrospinal fluid, such as antilymphocytes, antineuronal, antineurofilament, or antiglycolipid antibodies participate in the development of central nervous system lupus.

Many reports have implied an association between antihistone antibodies and various neurological diseases including central nervous system lupus, the myelopathy of lupoid sclerosis, and Degos’ disease, Guillain-Barré syndrome, migraine, chorea, and seizures. Using liposome or micelle purified antihistone antibodies, Misra et al and Yu et al showed that antihistone antibodies could react with and disturb the functions of lectin activated lymphocytes and neutrophil phagocytosis. As brain tissue contains a higher proportion of phospholipids than other organs, we speculate that antihistone antibodies may have a biological effect on nerve tissue and may derange its function once the autoantibodies pass into the brain. In this study, antihistone antibodies were purified from serum samples from patients with SLE and the effects of these antibodies on normal rat brain astrocytes and their binding capacity with brain tissue were investigated.

Materials and methods
PURIFICATION OF ANTICARDIOLIPIN ANTIBODIES WITH CARDIOLIPIN MICELLES
The purification of antihistone antibodies followed the method described by Harris et al with some modifications.

SUBCULTURE OF NORMAL RAT BRAIN ASTROCYTES
The RBA-1 cells were established by continuous passaging and enriching of cultures of the brain cells of three day old rats (JAR-2, F-51) for four years. The cells were cultured in 10% fetal bovine serum in RPMI 1640 without antibiotics. The RBA-1 cells were identified by the following properties: (a) no tumourigenicity when inoculated into nude mice; (b) 71-5% of the cells maintained diploid chromosomes in karyotype analysis; (c) the presence of glial fibrillary acidic protein in the cytoplasm detected by an immunochemical method; and (d) contact inhibition observed during long term culture.

[3H]Thymidine Incorporation of RBA-1 Cells
RBA-1 cells (100 μl; 1 × 10⁵/ml) were placed in flat bottomed microwells in triplicate. Antihistone antibodies (100 μl of 100, 200, and 400 μg/ml solutions) were added to the wells. In the control wells, 0-1 ml RPMI 1640 or the
corresponding concentration of human γ globulin (Cohn fraction II, Sigma Chemical, St Louis, MO, USA) was added instead of anticardiolipin antibodies. The mixture was incubated at 37°C in 5% carbon dioxide/95% air for 68 hours followed by four hours of pulsing with 18:5 kBq methyl \[^{3}H\]thymidine (specific activity 2.5×10^5 MBq/mmol; NEN, Boston, MA, USA). The cells were harvested and the radioactivity measured by a β counter.

ESTIMATION OF MEMBRANE POTENTIAL OF INDIVIDUAL RBA-1 CELLS

The membrane potential was measured by the method of Shapiro et al.26 The indicator dye used was 3, 3'-dihexyloxacarbocyanine iodide (Eastman Kodak, Rochester, NY, USA). The RBA-1 cells (5×10^5/ml) were cultured and reacted with 100 µg/ml anticardiolipin antibodies (or 100 µg/ml of γ globulin as control) in plastic petri dishes for one day before harvesting. The attached cells were collected by incubation with 0-125% trypsin-l mM EDTA in phosphate buffered saline solution (pH 7:2) for five minutes and the membrane potentials of these cells were estimated by EPICS C flow cytometry (Coulter Electronics, Hialeah, Miami, FL, USA) with excitation at 488 nm.

BINDING OF RBA-1 CELLS WITH ANTICARDIOLIPIN ANTIBODIES

RBA-1 (1×10^5 cells/0.1 ml) were reacted with 0·1 ml anticardiolipin antibodies (400 µg/ml) or human γ globulin (400 µg/ml) in an ice bath for 30 minutes. After two washes with Hanks’s balanced salt solution the cells were stained with FITC conjugated goat antihuman IgG in an ice bath for another 30 minutes. After washing with Isoton II (Coulter), the percentage and fluorescence intensity of RBA-1 cells were measured by EPICS C flow cytometry.

DETECTION OF GLIAL FIBRILLARY ACIDIC PROTEIN IN RBA-1 CELLS

The detection of glial fibrillary acidic protein was carried out by the indirect immunofluorescence method reported by Jou and Akimoto24 with some modifications. RBA-1 cells (1·5×10^5/ml) were cultured on coverslips and became attached after overnight incubation. Anticardiolipin antibodies, RPMI, or γ globulin were then added to the individual cultures. After 18 hours incubation the cells were fixed in 70% ethanol (at 4°C) for one hour and coated with 0·5% bovine serum albumin to reduce the non-specific binding with proteins. The coverslips were then incubated in a moist chamber with mouse monoclonal antibodies to glial fibrillary acidic protein (1:200 dilution in phosphate buffered saline) (BioMakor, Israel) for one hour. After rinsing with phosphate buffered saline, the coverslips were stained with FITC conjugated goat antimouse IgG (Jackson Immunoresearch Laboratories, West Grove, PA, USA) at 1:20 dilution for 30 minutes and rinsed again with phosphate buffered saline. The cells were examined with a fluorescence microscope.

IMMUNOFLOURESCENCE STAINING OF MOUSE BRAIN TISSUE WITH ANTICARDIOLIPIN ANTIBODIES AND ANTIBODIES TO GLIAL FIBRILLARY ACIDIC PROTEIN

BALB/c mouse brains were cut cryogenically and stained by indirect immunofluorescence. Briefly, 5 µm thick cryostat sections of brain tissue were first incubated with anticardiolipin antibodies (100 µg/ml), or 1:200 diluted mouse antibodies to glial fibrillary acidic protein, for one hour at room temperature. Human γ globulin (100 µg/ml) was the control protein. After washing with phosphate buffered saline, FITC conjugated antihuman IgG (or FITC conjugated antimouse IgG for glial fibrillary acidic protein stain) was added and incubated for another 30 minutes. The slides were rinsed, mounted, and examined under a fluorescence microscope.

STATISTICAL ANALYSIS

The results are given as mean (SD) for the whole study. Statistic significance was analysed by the non-parametric Wilcoxon signed rank test.

**Results**

EFFECT OF ANTICARDIOLIPIN ANTIBODIES ON THE GROWTH OF RBA-1 CELLS

The proliferation of RBA-1 cells was evaluated by the incorporation of \[^{3}H\]thymidine into the cells. Figure 1 shows that the inhibition of proliferation of RBA-1 cells by anticardiolipin antibodies was dose dependent and greater than that by human γ globulin. Kinetic studies showed that anticardiolipin antibodies (100 µg/ml) inhibited the proliferation of RBA-1 cells (20·6 (5·1)% of the control) maximally after 24 hours of interaction whereas human γ globulin had no significant effect on these cells (fig 2).

EFFECT OF ANTICARDIOLIPIN ANTIBODIES ON THE EXPRESSION OF GLIAL FIBRILLARY ACIDIC PROTEIN IN RBA-1 CELLS

The specific marker for differentiated astrocytes

![Graph](http://ard.bmj.com/)  

*γ Globulin  
*aCL

Figure 1 Dose dependent suppression of anticardiolipin antibodies (aCL) (○) and human γ globulin (■) on the \[^{3}H\]thymidine incorporation of RBA-1 cells. *p<0.05; **p<0.001.
is glial fibrillary acidic protein in the cytoplasm. To determine if anticardiolipin antibodies interfere with the expression of glial fibrillary acidic protein, the immunofluorescence staining of glial fibrillary acidic protein in RBA-1 cells was evaluated after two days of incubation with anticardiolipin antibodies or γ-globulin. Figure 3 shows that in the presence of anticardiolipin antibodies (100 μg/ml) the cells underwent conspicuous morphological changes, but the concentration of glial fibrillary acidic protein in the cytoplasm decreased, as shown by the faint fluorescence in the cells (fig 4). Unequal expression of cytoplasmic glial fibrillary acidic protein in the individual astrocytes and the presence of cells with scanty dendrites in fig 4 give the small differences in cytoplasmic fluorescence. Whether the anticardiolipin antibodies are suppressing glial fibrillary acidic protein in RBA-1 cells needs further investigation.

EFFECTS OF ANTICARDIOLIPIN ANTIBODIES ON THE MEMBRANE POTENTIAL OF INDIVIDUAL RBA-1 CELLS

To test whether the inhibitory effects of the anticardiolipin antibodies on RBA-1 cells were mediated through their membrane effect, the membrane potential of these cells was evaluated by flow cytometry after incubation with the antibodies for various times. Depolarisation of

![Figure 2](http://ard.bmj.com/)

**Figure 2.** Comparison of proliferation inhibition of RBA-1 by 100 μg/ml anticardiolipin antibodies (aCL) and human γ globulin after incubation for one, three, and five days.

![Figure 3](http://ard.bmj.com/)

**Figure 3.** Morphological changes of RBA-1 cells after incubation with 100 μg/ml human γ globulin (A) and 100 μg/ml anticardiolipin antibodies (B) for one day (original magnification x400). The cells firmly attached to the glass surface with multiple extended dendrites (A). These cells became rounded and the dendrites decreased or even disappeared in some cells after incubation with anticardiolipin antibodies (B).

![Figure 4](http://ard.bmj.com/)

**Figure 4.** Changes in fluorescence intensity of glial fibrillary acidic protein in the cytoplasm of RBA-1 cells after incubation with 100 μg/ml human γ globulin (A) or anticardiolipin antibodies (B) for one day (original magnification x400). The RBA-1 cells were cultured in coverslips, fixed in 70% ethanol, and stained with mouse monoclonal antibodies to glial fibrillary acidic protein and FITC conjugated goat antimouse IgG antibodies.

![Figure 5](http://ard.bmj.com/)

**Figure 5.** Changes in membrane potential of RBA-1 cells measured by flow cytometry after incubation with different reagents for two days. (A) RBA-1 incubated with medium: positive cells 67-11%, mean fluorescence intensity 110-77. (B) RBA-1 incubated with 100 μg/ml human γ globulin: positive cells 60-47%, mean fluorescence intensity 102-85. (C) RBA-1 incubated with 100 μg/ml anticardiolipin antibodies: positive cells 26-22%, mean fluorescence intensity 63-19.
Figure 7

The RBA-1 cells was noted in the presence of anticardiolipin antibodies (100 μg/ml) in one and two day cultures, whereas no distinct change could be found in the culture with human γ globulin and RPMI 1640 (fig 5). In another experiment, the binding of anticardiolipin antibodies (100 μg/ml) or γ globulin (100 μg/ml) with RBA-1 cells was measured by flow cytometry. The table shows that the percentage binding of RBA-1 with anticardiolipin antibodies was higher than that with γ globulin.

IMMUNOFLUORESCENCE STAINING OF MOUSE BRAIN BY ANTICARDIOLIPIN ANTIBODIES AND ANTIBODIES TO GLIAL FIBRILLARY ACIDIC PROTEIN

As brain tissue contains many phospholipids, immunofluorescence staining of BALB/c mouse brain sections with anticardiolipin antibodies was carried out to see whether anticardiolipin antibodies are capable of binding with these phospholipid epitopes. Figure 6 shows that positive immunofluorescence staining of mouse brain by anticardiolipin antibodies was seen in the corpus callosum, a cellular zone near the corpus callosum, and nerve cells scattered in other portions of the brain. In contrast, negligible fluorescence was found in the stain with human γ globulin (fig 7). In the immunofluorescence staining of glial fibrillary acidic protein, positive results were also found in the corpus callosum and other areas of the brain (data not shown). In contrast, cryosections of mouse liver were not stained by anticardiolipin antibodies or antibodies to glial fibrillary acidic protein (data not shown) with the same method, indicating the high affinity of anticardiolipin antibodies with the brain tissue.

Discussion

Anticardiolipin antibodies are found in 22–70% of patients with SLE with neurological disorders, including focal cerebral and ocular ischaemia, transverse myelopathy, peripheral neuropathy, and movement and seizure disorders. In most instances these disorders are thought to be the result of thrombosis or vasculopathy induced by immune complexes. Several workers have shown, however, that membrane reactive autoantibodies such as antineuronal antibodies are correlated with neuropsychiatric disorders of SLE. Bluestein et al. found that integrins and heat shock proteins (hsp90) were potentially important membrane target mol-
Effects of anticardiolipin antibodies on astrocytes

cules for antineuronal and lymphocytotoxic antibodies in patients with neuropsychiatric lupus.

In addition to these observations there is some evidence that anticardiolipin antibodies also react with many types of activated cells in cell membranes. Khamashina et al.9 showed that purified anticardiolipin antibodies bound to phosphatidylethanolamine, phosphatidylserine, and phosphatidylinositol epitopes expressed on the freeze thawed platelets and red blood cells. Schorer et al.30 found that liposome purified anticardiolipin antibodies selectively inhibited prostacyclin release from thrombin activated endothelial cells. Misra et al.31 and Yu et al.32 showed that affinity purified anticardiolipin antibodies suppressed the proliferation of phytohaemagglutinin activated peripheral lymphocytes and phagocytosis of neutrophils. Dostal-Johnson et al.33 found that serum samples from patients with lupus anticoagulant contained antibodies against phosphatidylserine, cardiolipin, phosphatidylinositol, phosphatidylglycerol, and phosphatidylethanolamine. Brain tissue contains a high proportion of phospholipids and may become the target of anticardiolipin antibodies, especially in patients with SLE.

It is possible that the blood-brain barrier is constructed by microvascular endothelial cells and astrocytes.32 Functionally, this structure prohibits noxious substances such as autoantibodies from entering brain tissue. Although Lolli et al.34 found that the local synthesis of anticardiolipin antibodies within the central nervous system was possible in patients with multiple sclerosis, neurophilis, or Guillain-Barré syndrome, a simultaneous occurrence of anticardiolipin antibodies in the cerebrospinal fluid and serum of patients with SLE suggests that anticardiolipin antibodies may diffuse from serum to the central nervous system through a damaged blood-brain barrier.35

In an attempt to further understand the pathological part played by anticardiolipin antibodies on the central nervous lesions of SLE, normal rat astrocytes were used as target cells. We showed that micelle purified anticardiolipin antibodies in serum samples from patients with SLE exerted suppressive effects on these cells in a dose dependent manner. This observation has not been reported elsewhere. The inhibitory effect of anticardiolipin antibodies on normal rat brain astrocytes is important as about 75% of patients with neuropsychiatric SLE have blood-brain barrier damage.36,37 When RBA-1 cells were incubated with anticardiolipin antibodies, the astrocytes became spherical in appearance and the cell membrane depolarised. It appears that anticardiolipin antibodies derange the structure and function of the blood-brain barrier, probably through their membrane effect.21 Once the blood-brain barrier is damaged, noxious substances, including cytotoxic antibodies, can access the brain tissue causing neuropsychiatric disorders. In addition, direct binding of anticardiolipin antibodies to the cell membranes of RBA-1 cells, though only slight (table), was also seen in this study. This binding may interfere with the signal transduction of the cells. The epitope(s) on the surface membrane of RBA-1 cells for anticardiolipin antibody binding and the real mechanism for the interference of signal transduction after autoantibody binding were not investigated in this work. Another interesting observation in this study is that anticardiolipin antibodies bound to brain tissue but not to liver tissue. It seems that anticardiolipin antibodies selectively attack brain tissue and elicit neuropsychiatric lesions in SLE in addition to non-specific thrombus formation in the blood vessels. The tissue specific property of other autoantibodies in SLE has been shown by the selective renal damage caused by antibodies to double stranded DNA.36 Although we did not identify the target cells of the anticardiolipin antibodies in the brain, we noted that the nerve fibres and possibly glial cells in the corpus callosum, nerve cells in the cellular zone near the corpus callosum, and sporadic nerve cells in other brain areas were all positively stained by anticardiolipin antibodies (fig 6). Whether anticardiolipin antibodies cross react with antineuronal, antilymphocyte, anticytotoxic, and antineurifilament antibodies needs further investigation.

In conclusion, we found that anticardiolipin antibodies exerted inhibitory effects on normal rat astrocytes (blood-brain barrier damage) and bound directly to nerve fibres and nerve cells in the brain. This dual effect of anticardiolipin antibodies on astrocytes and neurons by anticardiolipin antibodies as autoantibodies to neural membranes play an important part in the pathogenesis of neuropsychiatric SLE.

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