Close association between valvar heart disease and central nervous system manifestations in the antiphospholipid syndrome

I Krause, S Lev, A Fraser, M Blank, M Lorber, L Stojanovich, J Rovensky, J Chapman, Y Shoenfeld

Background: Heart valves lesions and central nervous system involvement are among the most common manifestations of the antiphospholipid syndrome (APS).

Objective: To evaluate possible interrelations between these manifestations in a large group of APS patients.

Methods: 284 APS patients were evaluated retrospectively, 159 of whom had primary APS. Cardiac–CNS associations were determined for the entire study population, and for subgroups of patients with primary APS or APS associated with systemic lupus erythematosus (SLE).

Results: Significant associations where found between cardiac vegetations and epilepsy (p < 0.02), and between cardiac valve thickening or dysfunction and migraine (p = 0.002). Borderline association was found between valvar vegetations and migraine (p = 0.09). A significant association was also found between all valvar lesions and stroke or transient ischaemic attacks. Subanalyses showed that patients with primary APS had significant associations between cardiac valve pathology and all CNS manifestations, while patients with APS associated with SLE had no such associations.

Conclusions: The study suggests potential differences in biological behaviour between primary APS and APS associated with SLE. The presence of cardiac valve pathology may be a risk factor for several types of CNS involvement in PAPS.

Methods

Patients

We recruited 284 APS patients from seven medical centres in Israel, Yugoslavia, and the Slovak Republic. All patients met the 1997 revised Sapporo criteria for the antiphospholipid syndrome. All underwent detailed medical interview and routine physical examination by a qualified internist or rheumatologist before inclusion in the study. Further data were obtained from the patients’ medical files with respect to expression of the clinical manifestations of the disease, demographic variables, and laboratory results. The study was designed to evaluate possible associations between cardiac valve abnormalities and epilepsy or migraine. All patients underwent cross sectional and Doppler colour flow echocardiographic examination for determination of cardiac valve

Abbreviations: APL, antiphospholipid antibodies; APS, antiphospholipid syndrome; PAPS, primary antiphospholipid syndrome
abnormalities. Valve thickening was defined as focal areas of increased echogenicity and thickening of cardiac valve leaflets. For the mitral valve, a thickness of more than 5 mm was considered abnormal; for other valves abnormal thickness was determined according to standards of the specific medical centre. Valvar dysfunction was defined as the presence of stenosis or regurgitation of at least moderate severity. Non-infectious vegetations were defined as echocardiographically identified masses protruding from valve leaflets in the presence of sterile blood cultures and the absence of fever. Epilepsy was defined as at least two separate episodes of unprovoked seizures with an interval greater than 24 hours. Seizures could either be simple partial, complex partial, or generalised tonic–clonic, and had to be unrelated to metabolic abnormalities or to drug use or withdrawal, and without other provoking factors such as hypoxia or cerebral hypoperfusion. Migraine was defined by a neurologist at the specific medical centre as a recurrent severe unilateral or bilateral headache, lasting 4–72 hours, with or without aura, after excluding other relevant aetiologies such as tension headache or headache secondary to hypertension.

The cardiac–CNS associations were determined for the entire study population, and for subgroups of patients with primary APS (PAPS) or APS associated with systemic lupus erythematosus (SLE).

### Detection of anti-β2GPI antibodies and anticardiolipin antibodies

The levels of antiphospholipid antibodies were detected by enzyme linked immunosorbent assay (ELISA). Ninety six well ELISA plates (Nunc, Kamstrup, Roskilde, Denmark) were coated with cardiolipin (Sigma, St Louis, Missouri, USA), 50 mg/ml in ethanol, or β2 glycoprotein I (β2GPI), 10 μg/ml in phosphate buffered saline (PBS). Following blocking with 3% bovine serum albumin (BSA), sera at different dilutions were added and incubated for two hours at room temperature. Bound antibodies were detected using goat antihuman IgG or IgM conjugated to alkaline phosphatase (Sigma) and appropriate substrate. The colour reaction was read in a Titertrek ELISA reader (SLT Labistruments, Salzburg, Austria) at an optical density of 405 nm. Extensive washing with PBS followed each step.

### Statistics

Either the χ² test or Fisher’s exact test was applied as appropriate to analyse statistically significant differences between categorical variables. Two sided probability (p) values of less than 0.05 were considered significant.

### RESULTS

We studied 284 APS patients, 238 of whom (83.8%) were women and 46 (16.2%) were men. The mean (SD) age of the population was 42.5 (14.6) years. The most frequent clinical manifestations in the study population are shown in Table 1. The most common cardiac valve abnormalities were in patients with antiphospholipid syndrome with or without CNS manifestations (epilepsy and migraine) as shown in Table 2.

### Table 1 Clinical manifestations in the study population

<table>
<thead>
<tr>
<th>Clinical manifestations</th>
<th>Entire APS population (n = 284)</th>
<th>PAPS (n = 159)</th>
<th>SLE with APS (n = 97)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Deep venous thrombosis</td>
<td>57.4</td>
<td>52.2</td>
<td>59.8</td>
<td>NS</td>
</tr>
<tr>
<td>Arterial thrombosis</td>
<td>48.6</td>
<td>45.3</td>
<td>56.7</td>
<td>NS</td>
</tr>
<tr>
<td>Recurrent pregnancy loss</td>
<td>27.7</td>
<td>34.6</td>
<td>19.8</td>
<td>&lt;0.03</td>
</tr>
<tr>
<td>TIA/CVA</td>
<td>35.2</td>
<td>35.8</td>
<td>39.2</td>
<td>NS</td>
</tr>
<tr>
<td>Myocardial infarction</td>
<td>7.4</td>
<td>5.7</td>
<td>10.3</td>
<td>NS</td>
</tr>
<tr>
<td>Epilepsy</td>
<td>9.9</td>
<td>6.9</td>
<td>15.5</td>
<td>&lt;0.03</td>
</tr>
<tr>
<td>Migraine</td>
<td>18.0</td>
<td>20.8</td>
<td>14.4</td>
<td>NS</td>
</tr>
<tr>
<td>Cardiac valve vegetations</td>
<td>6.0</td>
<td>4.4</td>
<td>9.3</td>
<td>NS</td>
</tr>
<tr>
<td>Cardiac valve thickening</td>
<td>18.0</td>
<td>17.6</td>
<td>19.6</td>
<td>NS</td>
</tr>
<tr>
<td>Overall cardiac valve abnormalities</td>
<td>21.1</td>
<td>18.2</td>
<td>27.8</td>
<td>NS</td>
</tr>
<tr>
<td>Livedo reticularis</td>
<td>15.8</td>
<td>11.3</td>
<td>25.8</td>
<td>&lt;0.005</td>
</tr>
<tr>
<td>Thrombocytopenia</td>
<td>28.5</td>
<td>22.6</td>
<td>40.2</td>
<td>&lt;0.005</td>
</tr>
<tr>
<td>Autoimmune haemolytic anaemia</td>
<td>10.2</td>
<td>7.5</td>
<td>17.5</td>
<td>&lt;0.02</td>
</tr>
</tbody>
</table>

Values are per cent.

APS, antiphospholipid syndrome; CVA, cerebrovascular accident; PAPS, primary antiphospholipid syndrome; SLE, systemic lupus erythematosus; TIA, transient ischaemic attack.

### Table 2 Cardiac valve abnormalities in patients with antiphospholipid syndrome with or without CNS manifestations (epilepsy and migraine)

<table>
<thead>
<tr>
<th></th>
<th>With epilepsy</th>
<th>Without epilepsy</th>
<th>p Value</th>
<th>With migraine</th>
<th>Without migraine</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>(A) Whole study population</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiac valve vegetations</td>
<td>17.9</td>
<td>4.7</td>
<td>&lt;0.02</td>
<td>11.8</td>
<td>4.7</td>
<td>0.09</td>
</tr>
<tr>
<td>Valvar thickening and/or dysfunction</td>
<td>25.0</td>
<td>17.1</td>
<td>NS</td>
<td>33.3</td>
<td>14.6</td>
<td>0.002</td>
</tr>
<tr>
<td>Overall cardiac valve abnormalities</td>
<td>32.1</td>
<td>19.9</td>
<td>NS</td>
<td>37.3</td>
<td>17.6</td>
<td>0.002</td>
</tr>
<tr>
<td>(B) PAPS</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiac valve vegetations</td>
<td>18.1</td>
<td>3.5</td>
<td>&lt;0.03</td>
<td>12.1</td>
<td>2.4</td>
<td>&lt;0.04</td>
</tr>
<tr>
<td>Valvar thickening and/or dysfunction</td>
<td>27.3</td>
<td>16.9</td>
<td>NS</td>
<td>33.3</td>
<td>13.5</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Overall cardiac valve abnormalities</td>
<td>27.3</td>
<td>17.6</td>
<td>NS</td>
<td>36.3</td>
<td>13.5</td>
<td>0.002</td>
</tr>
<tr>
<td>(C) APS associated with SLE</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiac valve vegetations</td>
<td>13.3</td>
<td>8.5</td>
<td>NS</td>
<td>7.1</td>
<td>9.6</td>
<td>NS</td>
</tr>
<tr>
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<td>20.0</td>
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<td>NS</td>
</tr>
<tr>
<td>Overall cardiac valve abnormalities</td>
<td>33.3</td>
<td>26.8</td>
<td>NS</td>
<td>28.6</td>
<td>27.7</td>
<td>NS</td>
</tr>
</tbody>
</table>

Values are per cent.

(A): the whole study population (n = 284); (B): patients with PAPS (n = 159); (C): patients with SLE and APS (n = 97).

APS, antiphospholipid syndrome; PAPS, primary antiphospholipid syndrome; SLE, systemic lupus erythematosus.
Table 3  Cardiac valve abnormalities in patients with antiphospholipid syndrome with and without cerebrovascular accident or transient ischaemic attack

<table>
<thead>
<tr>
<th></th>
<th>With CVA/TIA</th>
<th>Without CVA/TIA</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>(A) Whole study population</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiac valve vegetations</td>
<td>10.0</td>
<td>3.8</td>
<td>&lt;0.04</td>
</tr>
<tr>
<td>Valvar thickening and/or dysfunction</td>
<td>26.0</td>
<td>13.6</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Overall cardiac valve abnormalities</td>
<td>32.0</td>
<td>15.2</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>(B) PAPS</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiac valve vegetations</td>
<td>7.0</td>
<td>2.9</td>
<td>NS</td>
</tr>
<tr>
<td>Valvar thickening and/or dysfunction</td>
<td>26.3</td>
<td>12.7</td>
<td>0.03</td>
</tr>
<tr>
<td>Overall cardiac valve abnormalities</td>
<td>28.1</td>
<td>12.7</td>
<td>&lt;0.02</td>
</tr>
<tr>
<td>(C) APS associated with SLE</td>
<td></td>
<td></td>
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DISCUSSION

In this study we have identified significant associations between cardiac valve lesions and CNS manifestations in a large group of APS patients. The presence of valvar vegetations was found to be associated with epilepsy and migraine, while valvar thickening or dysfunction, as well as overall valvar abnormalities, were significantly associated with the presence of migraine. Furthermore, cardiac valve vegetations or dysfunction, or both, were strongly associated with a high rate of CVA or TIA. The rate of cardiac valve abnormalities in our study population was 21.1%. It was somewhat more common in SLE patients (27.8%) than in primary APS (18.2%, NS). The prevalence of valvar pathology in our study is in the range reported in previous studies, ranging usually between 32% and 38% of APS patients, although lower and higher prevalences have also been reported. Recently, Zavaleta et al. reported a 70.8% rate of valvar lesions in APS patients, detected by transoesophageal echocardiography. New valvar lesions or progression of existing lesions were detected in nine of 12 patients during a five year follow up. Epilepsy was diagnosed in 9.9% of the patients in our study, and migraine in 18.0%—in line with previously reported rates. In the current study we did not have information on the sequence of events. Thus, further prospective studies are needed to evaluate whether cardiac valve disease is a risk factor for CNS manifestations, or vice versa.

A possible link between heart pathology and various CNS involvements in APS has not yet been investigated. The only cardiac–CNS association in APS to be reported thus far has been on valve vegetations as a possible risk factor for cerebral emboli. The results of our study support this observation. To the best of our knowledge a relation between valvar vegetations or dysfunction and other CNS manifestations has not yet been reported.

The pathogenesis of cardiac valve and cerebral abnormalities in APS is not entirely clear. It has been postulated that aPL cause valvar or endothelial injury directly, independently of the severity of the disease, with aPL deposition on the valves resembling immune complex deposition in the dermo-epidermal junction or in the kidney basement membrane in patients with SLE. Antibody deposition and complement components initiating valve damage have been described, along with increased endothelial cell expression of α3β1 integrin. On the other hand, some histological studies suggest that fibrin deposits are the major finding, not inflammation. It has also been suggested that antigenaemia may improve cardiac valvar lesions, though other studies did not show benefit of anticoagulant or antiaggregate therapy. In our study we did not find a relation between anticoagulant/antiaggregate treatment and the rates of valvar pathology.

As to CNS involvement, the specific role of aPL remains one of the least understood aspects of this syndrome. The link between aPL and epilepsy has been supported by several studies, but there is as yet no direct evidence that aPL per se are pathogenic or are direct mediators in the development of thrombotic or other neurological complications in human patients. Laboratory findings which indicate that aPL directly bind to brain tissue, inhibit astrocytes proliferation in vitro,
or depolarise brain nerve terminals may provide a basis for understanding the pathogenesis of epilepsy and other neurological complications in APS.29 30 The pathophysiology of migraine is complex and includes genetic, metabolic, vascular, and emotional components. Cuadrado et al31 reported on five APS patients with intractable headaches treated with a seven day course of anticoagulation. All five patients had a marked improvement of the headaches following the treatment. It was speculated that low dose anticoagulant therapy might be tried in APS patients with migraine.32 Our study was not designed to evaluate the role of treatment on CNS or cardiac manifestations in APS, but we observed similar rates of migraine in patients treated or untreated with either anticoagulants or antiaggregate agents. The linkage between cardiac valve and CNS pathologies found in our study suggests a common pathogenic mechanism mediated, for example, by localised aPRL associated tissue damage. Further studies are required to evaluate these mechanisms.

Of special interest are the opposite cardiac–CNS relations found between patients with primary APS and those with APS associated with SLE. Whereas primary APS patients showed a significant association between cardiac valve pathology and CNS manifestations, such associations were not found in patients with APS associated with SLE. Possible disparities in clinical presentation between patients with primary and secondary APS have not yet been thoroughly investigated. In a retrospective study, Asherson et al reported on a lower female to male sex ratio and lower rates of livedo reticularis and chorea in primary APS than in patients with SLE.33 Vianna et al found similar clinical and laboratory profiles in patients with primary APS or APS associated with SLE, with the exception that autoimmune haemolytic anaemia, endocardial valve disease, neuropathy, and low C4 levels all occurred more often in SLE patients.34 The different valvoc–CNS connections found in our study may reflect distinct pathogenic mechanisms underlying cardiac or cerebral involvements in APS patients, depending on their background autoimmune condition.

In conclusion, we report on a close association between cardiac valve pathology and several CNS manifestations, such as epilepsy, migraine, and CVA/TIA, in patients with APS. Our study also suggests that there are distinct biological behaviours in primary APS and in APS associated with SLE. From our results, the presence of cardiac valve pathology may be a risk factor for several types of CNS involvement in primary APS.

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