Isolation of Chlamydyphila pneumoniae from atheromas of the carotid artery and their antibiotics susceptibility profile

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BACKGROUND. Atherosclerosis is pathogenically similar to a chronic inflammatory response. Peripheral arterial disease (PAD) is a common manifestation of atherosclerosis. Chlamydyphila pneumoniae has been suggested to play a role in the origin of PAD.

OBJECTIVE. To determine whether C. pneumoniae is present in atherosclerotic lesions of the carotid artery wall in patients with PAD through several diagnostic methods and to characterize C. pneumoniae susceptibility profiles.

METHODS. The presence of C. pneumoniae in 9 tissue samples from atherosclerotic lesions obtained by carotid endarterectomy was investigated by 3 methods. Karnofsky-fixed specimens were examined by transmission electron microscopy (TEM), isolation of C. pneumoniae was attempted in LLCMK2 cell culture (ICC), and the presence of chlamydial DNA was investigated by polymerase chain reaction (PCR). The in vitro activities of azithromycin, roxitromycin and penicillin were tested in 4 isolations and the reference strain of C. pneumoniae AR39.

RESULTS. C. pneumoniae was detected in atherosclerotic plaques from 4 patients with PAD. The pathogen was identified by TEM, PCR and ICC. We report data of the in vitro susceptibility of 4 strains. These strains did not differ from those of the reference strain of C. pneumoniae AR39.

CONCLUSIONS. The profiles of antibiotic susceptibility of C. pneumoniae isolated from 4 of the patients did not differ from those of the reference strain.

Key words: C. pneumoniae. Atherosclerosis. Antibiotic susceptibility.

Aislamiento de Chlamydyphila pneumoniae de ateromas de la arteria carótida y su perfil de sensibilidad a los antimicrobianos

ANTECEDENTES. La patogénesis de la aterosclerosis es similar a una respuesta inflamatoria crónica. La enfermedad arterial obstructiva periférica (EAOP) es una manifestación común de la aterosclerosis. Se ha sugerido que Chlamydyphila pneumoniae participa en la EAOP.

OBJETIVO. Determinar, a través de varios métodos de diagnóstico, si C. pneumoniae está presente en lesiones ateroscleróticas de la arteria carótida de pacientes con EAOP. Posteriormente, estudiar la sensibilidad a los antibióticos de los aislamientos de C. pneumoniae.

MÉTODOS. La presencia de C. pneumoniae fue investigada a través de tres métodos en nueve lesiones ateroscleróticas carotídeas, obtenidas por endarterectomía. Las muestras fueron fijadas en la solución de Karnofsky para ser examinadas por microscopía electrónica; se intentó el aislamiento de C. pneumoniae en cultivos de células LLCMK2 (ACC); además, investigamos la presencia de ADN de clamidia por la reacción en cadena de la polimerasa (PCR). Se estudió in vitro la sensibilidad de los cuatro aislamientos y de la cepa de referencia de C. pneumoniae AR39 a los antimicrobianos azitromicina, roxitromicina y penicilina.

RESULTADOS. C. pneumoniae fue descubierta en cuatro placas ateroscleróticas de pacientes con EAOP. Fue identificada por microscopia electrónica, por PCR y también por ACC. Se obtuvieron los datos de la sensibilidad in vitro de los cuatro aislamientos. No se observó diferencia entre los perfiles de susceptibilidad a azitromicina, a roxitromicina y a penicilina, entre la cepa de referencia respiratoria de C. pneumoniae AR39 y los aislamientos vasculares.

CONCLUSIONES. La C. pneumoniae se encuentra frecuentemente en las lesiones ateroscleróticas avanzadas de las arterias carótidas de pacientes que sufren endarterectomías. Aunque estos hallazgos no establecen causalidad en la aterosclerosis de la arteria carótida, ellos deben estimular la investigación de un posible papel.
Atherosclerosis is pathogenically similar to a chronic inflammatory response. Peripheral arterial occlusive disease (PAOD) is a common manifestation of atherosclerosis. PAOD is defined as obstructive arterial disease of the lower extremities that reduces arterial flow during exercise or, in advanced stages, at rest. PAOD represents a marker for premature cardiovascular events (eg, myocardial infarction, stroke) and vascular-related death. Atherosclerosis is pathogenically similar to a chronic inflammatory response. Peripheral arterial occlusive disease (PAOD) is a common manifestation of atherosclerosis. PAOD is defined as obstructive arterial disease of the lower extremities that reduces arterial flow during exercise or, in advanced stages, at rest. PAOD represents a marker for premature cardiovascular events (eg, myocardial infarction, stroke) and vascular-related death.

Methods

Twelve patients who were having carotid endarterectomy were enrolled in this study, but only nine samples arrived at the laboratory in optimal conditions. All had advanced atherosclerotic plaques. The tissues studied were obtained from nine patients with PAOD during endarterectomy surgery between October 2001 and April 2002, at the Hospital Allende and Hospital Cordoba (Cordoba, Argentina). The local ethics committee approved the study protocol. Informed consent was obtained from all patients. The study complied with the Declaration of Helsinki. The patients who were admitted to our emergency room because of acute chest pain were diagnosed as having unstable angina pectoris and were included positive and negative controls, consisting in LLCMK2 cell cultures infected and mock-infected with a prototype strain of C. pneumoniae. The prototype strain of C. pneumoniae was the AR39 strain and was purchased from ATCC.

Isolation of C. pneumoniae in cell culture

Cell line: LLCMK2 was obtained from Unidad de Estudios de Clasiómicosis, Cátedra de Microbiología, Facultad de Farmacia y Biociencias, UBA, Argentina. Host cells were grown in Eagle’s minimal essential medium with non-essential amino acids and 2mM L-glutamine (EMEM) (Gibco), and 10% fetal calf serum (Gibco), at 35 °C in 5% CO2.

For isolation of C. pneumoniae in LLCMK2 cell cultures tissue samples were placed in 5 ml SPG medium and homogenized with ice for 2 min in a sterile tissue grinder. The homogenate was cleared by centrifugation, inoculated to 4 well monolayers in 86-well microtiter plates and cultured at 1800 rpm at 35 °C for 60 min, and then supernatants were replaced by chlamydial isolation medium consisting of EMMEM with 1 ug of cycloheximide per ml (Sigma Chemical Co). Two blind passages were made after the first inoculation of each specimen. To find out whether samples were infected with C. pneumoniae, we carried out immunofluorescence assays (IFA) with monoclonal antibodies against Chlamydia lipo polysaccharides and with monoclonal antibodies specific for C. pneumoniae ( Dako). After 72h incubation the monolayers of each passage were fixed with ice-cold methanol. Isolation was considered positive if at least three cells of the infected monolayer showed the typical inclusion of C. pneumoniae. All samples were inoculated by triplicates.

Molecular detection of C. pneumoniae

A 437 base pair fragment corresponding to the C. pneumoniae Pat restriction fragment described by Campbell et al.17 was amplified by PCR, using primers HL-1 (5’-GGTTCTCAATAGGGCCCTACT-3’) and HR-1 (5’-TCATAAATCTAGGTTGTTT-3’). The previously described PCR conditions were modified slightly as follows: clinical samples (0.2 ml of each homogenate) were pelleted by centrifugation (15,000 rpm, 30 min) and inoculated in protromosomes (200 pg/ml)-0.5% Tween at 55 °C for 60 min. Subsequently, the samples were boiled for 10 min. The PCR was carried out in 0.2 ml tubes containing PCR buffer (10 mM Tris pH 8.3, 50 mM KCl, 0.1% Triton 100), 2.5 mM deoxynucleoside triphosphate, 2.5 mM MgCl2, 0.5 mM (each) primer, and 2 U of Taq DNA polymerase (Lucigen, Brazil). Amplifications were carried out in a thermal cycle (Biometra, Göttingen, Germany). The conditions for PCR were as follows: reaction volumes were first heated for 2 min at 94 °C and then subjected to 35 amplification cycles of 1 min at 94 °C, 1 min at 55 °C, and 1 min at 72 °C. After the last cycle, samples were held for 7 min at 72 °C.

PCR products were visualized through 1.5% agarose gel stained with ethidium bromide. The sensitivity of the reaction was analyzed by serial dilutions of a known bacterial suspension and it was determined to be of 1 IU/reaction tube (not shown).

Negative samples were aliquoted and seeded with positive control (100 inclusion forming units (IFU) of DNA and was triplicated to run out an inhibition of the PCR.

Transmission electron microscopy

For transmission electron microscopy, the arterial tissue was processed by a standard technique, sectioned, and stained with lead citrate and uranyl acetate. Each specimen was examined with a Jeol 1200 EX (Jeol, Tokyo, Japan) transmission electron microscope. Ultra-thin sections were examined for bacterial structure: elementary and reticulate bodies (EB and RB) compatible with chlamydial organisms. Controls were processed in parallel with the samples from each patient.

Antibiotic susceptibility profiles

The antimicrobial agents azithromycin (Aventis Pharma, Bauniville, France), penicillin G (Sigma, USA), and amoxicillin (Pfizer, USA) were tested against the 12 patients. They were incubated in Karmarkov solution for TEM. Every assay in-
Searle, Supertal) were provided as powders and solubilized according to the instructions of the manufacturers. Susceptibility tests were performed on LLCMK₂ monolayers grown in 96-well plates as described previously. Briefly, after the cell monolayer reached confluence, the growing medium was replaced by serum-free Eagle’s minimal essential medium supplemented with 1 μg of cycloheximide per ml (Sigma, USA) and one of the studied antibiotics in twofold dilutions. Each well was inoculated with 0.1 ml of the test strain diluted to yield 10³ inclusion-forming units (IFU/ml) per ml. After 72 h incubation the monolayers were fixed and an IFA was performed as stated before.

The MIC (minimal inhibitory concentration) was defined as the lowest concentration at which no IFU were observed. In accordance with established protocols, the minimal chlamydial concentration (MCC) was subsequently determined by removing the drug-containing medium, washing the wells with phosphate-buffered saline, and adding fresh medium without antibiotics for further incubation for 72 h.

The cells were then disrupted with sterile glass beads and vigorous shaking for 2 h, harvested and inoculated on fresh monolayers, which were incubated for further 72 h, and stained as described above. The MCC was the lowest drug concentration that inhibited the production of IFU in the antibiotic-free passage. All experiments were made by quadruplicate.

Results

Inclusion-forming organisms which were IFA positive, were isolated from the carotid artery tissue samples B, H, I and J. Figure 1 shows photomicrographs of the ICC of sample B (fig. 1). PCR detection resulted equally sensitive for the four positive samples (fig. 2). We were able to detect amplification of strains B, H, I and J during the first 4 passages. The results are shown in table 1.

Structures similar in morphology and size to those of elementary (EB) and reticular bodies (RB) of C. pneumoniae were detected in four of the specimens when analyzed by TEM (fig. 3B). The EB were detected as particles with a diameter of 100 to 300 nm, surrounded by a membrane, and containing an electron-dense core measuring up to 100 nm in diameter (fig. 3B). The EB and RB were also identified in macrophages cells (fig. 3A).

The four samples positive for C. pneumoniae were classified as grade 3 lesions.

The four isolations positive from carotid lesion B, H, I, J and respiratory reference strain C. pneumoniae AR39 were compared for antibiotic susceptibility profiles. The results of MIC and MCC tests are summarized in table 2.

### Table 1. Patients and specimens examined

<table>
<thead>
<tr>
<th>Plaques</th>
<th>Macro grade</th>
<th>ICC</th>
<th>PCR</th>
<th>TEM</th>
<th>Age</th>
<th>Sex</th>
<th>Risk factors*</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>2</td>
<td>–</td>
<td>Negative</td>
<td>N**</td>
<td>52</td>
<td>M</td>
<td>HLB-HT</td>
</tr>
<tr>
<td>B</td>
<td>3</td>
<td>+</td>
<td>Positive</td>
<td>EB-RB***</td>
<td>56</td>
<td>M</td>
<td>HT-CS</td>
</tr>
<tr>
<td>C</td>
<td>3</td>
<td>–</td>
<td>Negative</td>
<td>N</td>
<td>74</td>
<td>F</td>
<td>D-HT</td>
</tr>
<tr>
<td>D</td>
<td>2</td>
<td>–</td>
<td>Negative</td>
<td>N</td>
<td>74</td>
<td>M</td>
<td>D-HT</td>
</tr>
<tr>
<td>F</td>
<td>2</td>
<td>–</td>
<td>Negative</td>
<td>N</td>
<td>56</td>
<td>F</td>
<td>HLB-HT</td>
</tr>
<tr>
<td>G</td>
<td>3</td>
<td>–</td>
<td>Negative</td>
<td>N</td>
<td>77</td>
<td>F</td>
<td>HLB-HT-CS-O</td>
</tr>
<tr>
<td>H</td>
<td>3</td>
<td>+</td>
<td>Positive</td>
<td>EB-RB</td>
<td>57</td>
<td>M</td>
<td>HLB-HT-CS-O</td>
</tr>
<tr>
<td>I</td>
<td>3</td>
<td>+</td>
<td>Positive</td>
<td>EB-RB</td>
<td>72</td>
<td>M</td>
<td>HT-CS</td>
</tr>
<tr>
<td>J</td>
<td>3</td>
<td>+</td>
<td>Positive</td>
<td>EB-RB</td>
<td>66</td>
<td>M</td>
<td>HLB-HT-CS-O</td>
</tr>
</tbody>
</table>

*Risk factors: HT: hypertension; CS: cigarette smoking; HLB: high lipids levels in the blood; D: diabetes mellitus; O: obesity
**N: None bacterial structure was observed.
***EB-RB: elemental bodies and reticular bodies.
There is evidence for a strong association between chronic infection with *C. pneumoniae* and smoking. It is concluded from the present data that chronic infection with the pathogen would not be an independent risk factor for carotid atheromatous lesions.

This study provides direct evidence to show the presence of viable *C. pneumoniae* in the carotid atheroma of patients with PAOD of Córdoba, Argentina. *C. pneumoniae* is a pathogenic organism which has been shown in vitro to be capable of infecting aortic smooth muscle cells, endothelial cells, and macrophages; all of which are involved in atherogenesis. The finding of the organism in the arterial lesion does not prove that *C. pneumoniae* is the leading cause in the progression of atherosclerosis. However, having a known pathogen replicating in the atheroma cells suggests that *C. pneumoniae* might be involved in the pathogenesis of PAOD.

Systemic dissemination of *C. pneumoniae* from the respiratory tract to the cells of the vascular wall requires a cellular transport system. Circulating monocytes, which are pivotal to the development of atherosclerosis, are known to migrate into the vascular wall. The finding of *C. pneumoniae* in macrophage cells raises the possibility that the bacteria reach the vascular wall through the infection of monocytes.

Numerous electron microscopy studies have more finely characterized the morphology of EB and RB, the continuum of intermediate forms, and the inclusion membrane within which the entire growth phase of the development takes place. In this study we observed structures similar to inclusions of *C. pneumoniae* and typical elementary bodies (pear-shaped) in the macrophage cells of original specimens from atheromatous plaque. Extensive work is needed to determine if the bacteria can complete an infective cycle in those cells.

This is the first study that describes the antibiotic susceptibility profile of peripheral arterial strains in Argentina. It is important to note that antibiotics have been demonstrated to be active in vitro against *C. pneumoniae*. Thus, chlamydial eradication from cells other than monocytes/macrophages is potentially feasible. However, there is evidence suggesting that persistence may be established in those cells, too.

The results of trials investigating a possible effect of antibiotics on coronary artery disease are contradictory. Negative outcomes might be explained not only by the use of an incorrect hypothesis (that infection is not atherogenic) but also by an inadequate sample size or by the use of an ineffective antibiotic regimen.

Non-selective use of roxithromycin is inadequate for prevention of restenosis after coronary stenting. In patients with high titres, roxithromycin reduced the rate of restenosis. A possible role of *C. pneumoniae* in peripheral arterial occlusive disease is intriguing but has rarely been investigated. A randomized trial of roxithromycin versus placebo was carried out with a subgroup of the patients. Antibiotic medication showed effective in inhibiting progression of PAOD.

A beneficial effect of 4-week roxithromycin treatment on PAOD was observed during a follow-up period of 2.7 years. In that study, *C. pneumoniae* seropositivity

**Minimal chlamydical concentration.**

**Minimal inhibition concentration.**

**Minimal chlamydial concentration.**

All MICs and MCCs values were within the range known for respiratory isolates, and all strains presented similar sensitivity profile to the antibiotics tested.

### Discussion

The first study that indicated a possible association between *C. pneumoniae* and coronary artery disease was performed in Finland and showed that patients with coronary artery disease were significantly more likely to have serologic evidence of past infection with *C. pneumoniae* than controls. Since that time, serologic studies from the United States and other countries have demonstrated similar findings among patients with coronary artery disease and in patients with thickening of the carotid arteries as well.

The important observations in this study are as follows: the detection of viable *C. pneumoniae* in 4/9 of arterial atheromatous lesions, in keeping with the results of some previous studies in which the organism has been detected in 71 to 100% of lesions. *C. pneumoniae* was detected in atherosclerotic plaques for all (44%) smoking patients with PAOD but was not detected in nonsmokers.

**TABLE 2. Activities of antibiotics against cardiovascular isolations (B, H, I and J) and respiratory strain AR29**

<table>
<thead>
<tr>
<th>Antibiotics</th>
<th>Concentration (mg/ml)</th>
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<tbody>
<tr>
<td></td>
<td>Strain AR29</td>
</tr>
<tr>
<td>Azithromycin</td>
<td>0.250 0.0625</td>
</tr>
<tr>
<td>Roxithromycin</td>
<td>0.125 0.0625</td>
</tr>
<tr>
<td>Penicillin</td>
<td>&gt; 2</td>
</tr>
</tbody>
</table>

**A** Transmission electron micrograph of a macrophage cell in an area of atheroma within an artery. The macrophage is identified as a cell full of lipid droplets and vacuoles containing *C. pneumoniae* (EB). The scale bar represents 1 μm. **B** Transmission electron micrograph of an atheromatous arterial wall. Elementary bodies of *C. pneumoniae* appear pear-shaped in some profiles (EB) and contain a central, electron-dense core. The scale bar represents 500 nm.

Figure 3. A) Transmission electron micrograph of a macrophage cell in an area of atheroma within an artery. The macrophage is identified as a cell full of lipid droplets and vacuoles containing *C. pneumoniae* (EB). The scale bar represents 1 μm. B) Transmission electron micrograph of an atheromatous arterial wall. Elementary bodies of *C. pneumoniae* appear pear-shaped in some profiles (EB) and contain a central, electron-dense core. The scale bar represents 500 nm.
had been associated with PAOD but not with coronary artery disease or cerebrovascular occlusive disease. A preference of *C. pneumoniae* for peripheral vessels is conceivable, since lower limb arteries have a different embryological origin and different histological constitution than coronary and cerebral arteries. That study provides strong evidence that *C. pneumoniae* is involved in the progression of PAOD and that antibiotic treatment directed against *C. pneumoniae* is effective in inhibiting this process.

Although *C. pneumoniae* infection can be effectively treated with tetracyclines and macrolide drugs, it is not known whether the organisms in atheroma could be eradicated by treatment or whether eradication of the organisms would have a favorable effect on the lesions. A controlled treatment trial with an effective antibiotic might be considered for future investigations. In our study, no difference was observed among the vascular isolates or between the isolates and the respiratory strains previously described. Apparently, vascular isolates of *C. pneumoniae* would not have a distinctive susceptibility profile. This confirms the finding that the *C. pneumoniae* isolates obtained from PAOD are very homogeneous in their genetic background, protein profile, and immunogenicity. However, since a genetic typing system has not yet been established for *C. pneumoniae*, the presence of a genotype with a distinct vascular pathogenicity cannot be excluded.

Our results emphasize the potential role of *C. pneumoniae* in the etiology of atherosclerosis and suggest that the development of atheroma could be prevented by antibiotic treatment in patients with proved *C. pneumoniae* infection.

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References