**Vascular Biology, Atherosclerosis, and Endothelium Biology**

**Chlamydophila pneumoniae Infection Induces Alterations in Vascular Contractile Responses**

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**Chlamydophila pneumoniae** infection has been associated in previous studies with coronary artery disease. The live bacterium has been detected within atherosclerotic plaques and can induce the structural remodeling of the vessel wall. However, the direct effects of infection on the contractile characteristics of the arteries remain unknown. Left anterior descending coronary arteries isolated from porcine hearts were dissected and placed in culture medium for 72 hours before infection with *C. pneumoniae*. Contractile responses to high molar KCl and u46619 levels and relaxation responses to bradykinin and sodium nitroprusside were assessed at days 5 and 10 postinfection. *C. pneumoniae* induced decreases in both KCl- and u46619-induced contractile responses at both time points. The altered contractile responses coincided with a down-regulation of L-type Ca2+ channels at both time points and inositol 1,4,5-triphosphate receptor (IP3R) levels at day 10 postinfection. A decrease in endothelial nitric oxide synthase expression (eNOS), the primary source of NO within the vessel wall, is up-regulated in response to *C. pneumoniae* infection. Endothelial dysfunction, a key initiating step in the progression of atherosclerosis, results primarily from a reduced bioavailability of nitric oxide (NO) leading to impaired endothelial-dependent relaxation. Infectious burden has been noted as an independent risk factor for endothelial dysfunction. Chronic and acute inoculations with the *C. pneumoniae* have been demonstrated to induce endothelial dysfunction in mice and pigs respectively. However, it is difficult to establish a causative role for *C. pneumoniae* in endothelial dysfunction with an in vivo system because of the involvement of an active immune response to infection. In vitro findings have provided some evidence for a direct role of the bacterium in promoting this dysfunctional state. *C. pneumoniae* infection can directly decrease endothelial nitric oxide synthase expression (eNOS), the primary source of NO within the vessel wall. Chlamydial heat shock protein 60 (chSP60), produced during active chlamydial replication, induces a decrease in eNOS expression and activity in ECs and stimulates endothelial dysfunction in isolated porcine coronary arteries. Endogenous HSP60, which is up-regulated in response to *C. pneumoniae* infection, promotes endothelial activation and is associated with endothelial dysfunction in adolescents.

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Whether \( C.\ pneumoniae \) infection can induce similar cellular changes within a complex vascular environment leading to endothelial dysfunction in the absence of an immune response is thus worth investigating.

VSMC proliferation, another key feature of atherosclerotic development, also could have important implications for the functional characteristics of arterial vessels. During the proliferative process, VSMCs undergo a phenotypic switch from a contractile (differentiated) VSMC to a synthetic (de-differentiated) VSMC. This de-differentiation process typically involves alterations in the expression of contractile proteins such as smooth muscle \( \alpha \)-actin and smooth muscle myosin heavy chain as well as \( \text{Ca}^{2+} \) handling proteins like the L-type \( \text{Ca}^{2+} \) channel.\(^{24}\)

As a result, the synthetic phenotype observed in proliferative VSMC could severely alter a vessel’s function and has been suggested to play an important pathophysiological role in the development of vascular diseases such as atherosclerosis, restenosis and hypertension.\(^{25}\) Direct \( C.\ pneumoniae \) infection and treatment with chSP60 can induce VSMCs proliferation:\(^{21,26,27}\) However, little is known about how \( C.\ pneumoniae \)-mediated proliferation can influence vascular function responses within a complete vessel system.

The current investigation aims to determine the direct action of \( C.\ pneumoniae \) infection on the functional responses of coronary arteries. We have recently developed an \( \text{ex vivo} \) porcine coronary artery model that allows us to determine the direct effect of \( C.\ pneumoniae \) infection within an arterial environment in the absence of an immune response.\(^{27}\) Using this approach, \( C.\ pneumoniae \) was able to infect the endothelial layer, migrate into the medial layer, and actively replicate within the vessel wall for up to 10 days postinfection. This active replication resulted in significant arterial thickening through induction of vascular smooth muscle cell (VSMC) proliferation. This current study uses this \( \text{ex vivo} \) coronary artery model to investigate vascular contractile and relaxation responses within \( C.\ pneumoniae \)-infected vessels before and after vascular hypertrophic responses.

### Materials and Methods

#### Reagents and Antibodies

Cycloheximide, bradykinin acetate, 9,11-Dideoxy-11\( \alpha \), 9\( \alpha \)-epoxy-methanoprostaglandin \( F_{2\alpha} \) (u46619), sodium nitroprusside, anti-\( \alpha \)-smooth muscle myosin, and anti-\( \alpha \)-smooth muscle actin antibodies were obtained from Sigma-Aldrich (St. Louis, MO). The anti-IP3 receptor-1 and -L-type of voltage-gated \( \text{Ca}^{2+} \) channel antibodies were obtained from Alomone Labs (Jerusalem, Israel). The anti-iNOS antibody was obtained from Abcam (Cambridge, MA). The eNOS antibody was obtained from BD Transduction Laboratories (Franklin Lakes, NJ). The anti-\( \text{NFkB} \) antibody was obtained from Santa Cruz (Santa Cruz, CA). The \( \text{IKB}\alpha \) antibody was obtained from Cell Signaling (Danvers, MA). The chlamydial HSP60 antibody was obtained from Affinity BioReagents (Golden, CO). The human HSP60 antibody was obtained from Stressgen (Ann Arbor, MI). The anti-NF\( \text{kB} \) antibody, HRP-conjugated anti-mouse IgG and HRP-conjugated anti-rabbit IgG antibodies were obtained from Millipore (Billerica, MA). The dihydroethidium, Alexa Fluor 488-conjugated anti-mouse IgG and Alexa Fluor 488-conjugated anti-rabbit IgG antibodies were obtained from Invitrogen (Carlsbad, CA).

#### Propagation of \( C.\ pneumoniae \)

\( C.\ pneumoniae \) AR39 strain was obtained from the University of Washington (Seattle, WA). The organism was propagated in HL cells as described previously.\(^{28}\) The purified organism was resuspended in chlamydial sucrose–phosphate–glutamate medium (SPG) and stored at −80°C until use. The titer of \( C.\ pneumoniae \) was determined in cycloheximide-treated HL cells, and concentrations used were expressed as inclusion-forming units (IFU) per milliliter.\(^{29}\)

#### Isolation and Infection of Porcine Coronary Arteries

Coronary artery isolation, culturing, and \( C.\ pneumoniae \) infection were performed as previously described.\(^{27}\) SPG and heat-inactivated \( C.\ pneumoniae \) (mock infection) were used for treatment and infection controls, respectively. After infection, vessels were maintained in culture for up to 10 days. Organ culture medium was changed (without antibiotics) every 48 hours, and the old medium was collected for conditioned medium experiments and protein analysis. \( C.\ pneumoniae \) infection induced arterial thickening in this model starting at day 6 postinfection. To determine whether alterations in vascular response preceded or followed hypertrophy, parameters were examined at two main time points, days 5 and 10 postinfection.

#### Vascular Function Assessment

Vascular function was assessed for coronary segments collected at days 4 and 5 and days 9 and 10 postinfection, using the experimental protocol previously described.\(^{27}\) All functional experiments were completed within a laminar flow biosafety cabinet to prevent spread of infection.

#### Treatment of Human Umbilical Vein Endothelial Cells with Conditioned Medium

Human umbilical vein endothelial cells (HUVECs) were obtained from Lonza and maintained in culture conditions with EGM-2 (Lonza, Walkersville, MD). Cells from P3 to P6 were used for experiments. HUVECs were plated at \( 2 \times 10^6 \) cells per well in a 6-well plate. Organ culture medium was collected every 48 hours during media changes. Media from similar treatments were combined in pairs; a portion (2 mL) of the medium was placed directly over HUVECS plated in 6-well plates. HUVECS were collected at 24 hours after incubation.
with conditioned medium. The additional medium (1 mL) was collected and spun at 1105 rpm for 5 minutes. The supernatant was collected and stored at −80°C for protein analysis.

**Western Blot Analysis**

To assess protein expression levels within treated vessels, coronary segments were collected at days 4 and 5 and days 9 and 10 postinfection. On collection, segments were rinsed with PBS, flash-frozen in liquid nitrogen and stored at −80°C. Coronary segments were subsequently ground and re-suspended in RIPA buffer (50 mmol/L Tris-HCl, pH 7.5, 150 mmol/L NaCl, 1 mmol/L EDTA, 1 mmol/L EGTA, 1% Triton 100, 0.1% SDS, 0.5% sodium deoxycholate, 1 µg/mL leupeptin, 1 mmol/L PMSF, 1 mmol/L protease inhibitor cocktail, 1 mmol/L DTT, and 1 mmol/L benzimidazole). HUVECs collected after conditioned medium treatment were re-suspended in RIPA buffer, sonicated, spun at 14,000 rpm for 10 minutes at 4°C, and the supernatant was collected for Western blotting. Protein assays were performed to determine concentrations within tissue and HUVEC lysates. Lanes were loaded with 5 µg of tissue lysate for detection of smooth muscle actin and myosin and 30 µg of protein for the other targeted proteins. Tissue samples were run in duplicate, with half of the blot used to probe with antibodies and the other half used for amido black staining. For HUVEC lysates, lanes were loaded with 30 µg of protein. Proteins were separated using a 4% to 12% gradient denaturing polyacrylamide gel and transferred electrophoretically onto a nitrocellulose membrane. Membranes were incubated with anti-smooth muscle actin (1:20,000), anti-smooth muscle myosin (1:1000), anti-iP3 receptor-1(1:500), anti-L-type of voltage-gated Ca2+ channel (1:500), anti-eNOS (1:200), anti-iNOS (1:500), anti-p22phox (1:500), anti-NF-κB (1:1000), anti-lkB (1:1000), or anti-GAPDH (1:5000) antibodies. HRP-conjugated anti-mouse IgG and anti-rabbit IgG were used as secondary antibodies. Bands were visualized with the Supersignal West Pico or Femto Chemiluminescent Substrates (Pierce). Bands and Amido Black staining were quantified by densitometry with Quantity One software (Bio-Rad).

In Situ Superoxide Detection and Immunofluorescence Staining

Coronary segments were collected at days 9 and 10 postinfection, fixed in ethanol, rehydrated, and subsequently equilibrated in 30% sucrose solution before mounting in OCT compound (Tissue-tek, Sakura Finetek, Torrance, CA). These OCT mounted segments were slowly frozen to −80°C, cut (HM 500 OM Cryostat, Microm) into 8-µm sections and placed on positively coated glass slides. To evaluate protein expression in coronary sections, antigen retrieval was first performed as previously described.27 Subsequently, sections were incubated with anti-eNOS(1:25), anti-p22phox (1:50), anti-TLR4 (1:10), anti-chSP60 (1:50) and anti-hSP60 antibodies. Texas Red–conjugated anti-mouse IgG (1:750), Alexa Fluor 488–conjugated anti-mouse IgG (1:1000), and Alexa Fluor 488–conjugated anti-rabbit IgG (1:1000) were used as secondary antibodies for the respective primary antibodies. Hoescht staining solution (5 ng/mL) was added to the slides to identify nuclei. To determine the levels of superoxide anions within the vessels, sections were incubated with dihydroethidium as previously described.30 Slides were fixed with Fluoro-Save reagent (Calbiochem, San Diego, CA) to preserve fluorescence. Staining was visualized using a fluorescent inverted microscope (TE-2000s, Nikon) and mean intensity was quantified using NIS Elements imaging software (Nikon, Mississauga, ON).

**Statistical Analysis**

Data are presented as mean ± SEM unless otherwise stated. Differences between treatment groups were assessed by one-way analysis of variance using the Student–Newman–Keuls method. A probability of P < 0.05 was considered statistically significant.

**Results**

*C. pneumoniae Induces Alterations in Vascular Contractile Function*

Isolated coronary vessels infected with *C. pneumoniae* displayed significant decreases in maximal contractile response to KCl when compared with control and mock-infected vessels at both days 5 (Figure 1A) and 10 postinfection (Figure 1B). Similar qualitative responses were used to construct a standard curve. Proteins were separated using a 9% gradient denaturing polyacrylamide gel and transferred electrophoretically onto a nitrocellulose membrane. Membranes were incubated with either anti-chSP60 (1:1000) or anti-hSP60 (1:1000) antibodies, and HRP-conjugated anti-mouse IgG was used as secondary antibody. Bands were visualized with the Super signal West Pico or Femto Chemiluminescent Substrates (Pierce). Bands and Amido Black staining were quantified by densitometry with Quantity One software (Bio-Rad).
found in response to receptor-mediated contraction with u46619 (Figure 1, C and D). At day 5 postinfection, the contractile response to both 150 and 300 nmol/L u46619 in C. pneumoniae–infected segments was decreased compared with control and mock-infected vessels. At day 10 postinfection, the contractile response to u46619 in infected segments was decreased only when compared with control.

C. pneumoniae Infection Alters Expression of Calcium-Handling Proteins but Not Contractile Proteins

Expression of the smooth muscle contractile proteins actin and myosin heavy chain was assessed in vessels at days 5 and 10 postinfection (Figure 2, A and B). Expression levels appeared to decrease with C. pneumoniae infection; however these changes were not significantly different compared with other treatment at day 5 (smooth muscle actin, \( P = 0.087; \) myosin, \( P = 0.172 \)) and day 10 (smooth muscle actin, \( P = 0.145; \) myosin, \( P = 0.143 \)). The expression of calcium handling proteins involved in smooth muscle contraction, L-type voltage gated Ca\(^{2+} \) channel and IP\(_3\) receptor, was also determined at days 5 and 10 postinfection (Figure 3, A and B). The expression of the L-type Ca\(^{2+} \) channel was significantly decreased in C. pneumoniae–infected vessels compared with control and mock-infected vessels at day 5 postinfection (Figure 3A) and day 10 (Figure 3B). The expression of IP\(_3\) receptor was unchanged between groups at day 5 postinfection (Figure 3C) but was significantly decreased in infected vessels compared with control at day 10 postinfection (Figure 3D).

C. pneumoniae Infection Attenuates Endothelial Dependent Relaxation in Isolated Coronary Vessels

Precontraction levels in response to 30 nmol/L u46619 remained unchanged between treatment groups at both days 5 and 10 postinfection (data not shown). Endothelial-dependent relaxation responses to bradykinin remained unaltered between groups at day 5 postinfection (Figure 4A), but at day 10 (Figure 4B) were significantly reduced in infected vessels compared with control at \( 10^{-9} \) to \( 10^{-6} \) mol/L and at \( 10^{-7} \) to \( 10^{-6} \) mol/L in comparison to mock-infected vessels (Figure 4B). No change in endothelial-independent relaxation responses to sodium nitroprusside was observed among groups at either time point (Figure 4, C and D).

C. pneumoniae Infection Induces Decrease in eNOS Expression

Expression of eNOS and iNOS, the nitric oxide producing enzymes within the vessel was determined via Western blot analysis. eNOS expression was unchanged among treatment groups at day 5 postinfection (Figure 5A). At day 10 postinfection, eNOS was down-regulated in infected vessels compared with control and mock-infected vessels (Figure 5B). Further analysis via immunostaining of coronary vessels at day 10 postinfection revealed a significant decrease in the mean fluorescence of infected...
cross-sections compared with control and mock-infected tissues (Figure 5C). iNOS expression remained unchanged between treatment groups at both time points (data not shown).

C. pneumoniae Components Stimulates Superoxide Anion Production and p22phox Expression

Expression of p22phox, a major subunit of NADPH oxidase, was determined via Western blot (Figure 6A) and immunostaining (Figure 6, B and C). Expression of p22phox, as detected by Western blots, was not changed among groups at day 5 postinfection but was up-regulated in both mock-infected and C. pneumoniae–infected vessels compared with control vessels at day 10 postinfection (Figure 6, A and B). p22phox immunostaining at day 10 postinfection confirmed these results, with higher fluorescence levels in mock-infected and C. pneumoniae–infected vessels compared with control (Figure 6C).

Superoxide anion production at day 10 postinfection was visualized using dihydroethidium staining on infected coronary cross-sections (Figure 7). Quantification of this staining revealed significant increases in mean fluorescence in both mock-infected and C. pneumoniae–infected vessels compared with control (a 99% (P < 0.05) and 207% (P < 0.01) change in comparison to control, respectively.

NFκB Activation Is Not Altered in Infected Coronary Arteries

The NFκB pathway is known to be involved in the inflammatory process within the vessel wall, as such both NFκB and IκB protein expression was assessed. Expression of both proteins did not significantly change between treatments at days 5 and 10 postinfection. The ratios of IκB to NFκB in both C. pneumoniae–infected and mock-infected vessels varied from control vessels but not enough to be statistically significant.

C. pneumoniae Increases TLR4, hHSP60, and cHSP60 Expression in the Endothelial Layer

TLR4, cHSP60, and hHSP60 localization along the endothelial layer was determined via immunostaining (Figure 8). Both C. pneumoniae–infected and mock-infected vessels displayed an increase in TLR4 and hHSP60 expression when compared with nontreated vessels (Figure 8, D and E compared with G and H). In addition, chSP60 expression could be localized along the endothelial layer in C. pneumoniae–infected vessels (Figure 8I) but was absent in both control and mock-infected coronary vessels.
Conditioned Medium Does Not Alter eNOS and p22phox Expression in HUVECs

Both eNOS and p22phox protein expression were evaluated in HUVECs treated with conditioned medium from days 4, 6, 8, and 10 postinfection via Western blot. Expression of both eNOS and p22phox in HUVECs treated with conditioned medium from all four time points were not significantly different between treatments (data not shown). cHSP60 and hHSP60 levels within conditioned medium from the different treatments were analyzed by Western blot. All conditioned medium displayed cHSP60 and hHSP60 levels below the bottom level of our standard curve of 0.01 μg of protein, equivalent to 0.08 μg/mL from the nonconcentrated conditioned medium (data not shown). cHSP60 could not be detected in culture medium from control and noninfected coronary vessels.

Figure 5. eNOS expression in infected isolated coronary arteries. A: Western blot analysis of whole tissue lysates at day 5 and day 10 postinfection. Representative Western blots are displayed above graph. Expression levels were represented as fold change over day 0 tissue ± SEM, n = 4 to 6. Representative IHC staining (B) quantification and (C) images of vessels collected at day 10 postinfection. Fluorescence intensity (B) is represented as mean ± SEM, n = 4. Hoescht, blue; eNOS, green. Scale bar = 100 μm. *P < 0.05 versus control and mock infection.

Figure 6. p22phox expression in infected isolated coronary arteries. A: Western blot analysis of whole tissue lysates at day 5 and day 10 postinfection. Representative Western blots are displayed above graph. Expression levels were represented as fold change over day 0 tissue ± SEM, n = 4 to 6. Representative IHC staining (B) quantification and (C) images of vessels collected at day 10 postinfection. Fluorescence intensity (B) is represented as mean ± SEM, n = 4. Hoescht, blue; p22phox, green. Scale bar = 100 μm. †P < 0.05 versus control.
Discussion

We hypothesized that C. pneumoniae infection may participate in phenotype switching in VSMCs and as such may interfere with the ability of these cells to properly mediate the contractile and relaxation responses within an arterial environment. Our findings clearly demonstrate that C. pneumoniae infection leads to an impairment of both receptor (u46619) and depolarization (KCl)–mediated contractile responses (Figure 1). The involvement of the L-type Ca\(^{2+}\) channel and IP3R in this impairment was evaluated in the present study. In VSMCs, the L-type Ca\(^{2+}\) channel is the most important ion channel for the influx of extracellular Ca\(^{2+}\) leading to downstream activation of myosin light chain kinase (MLCK). The channel can be opened directly through membrane depolarization by high concentrations of K\(^+\) or indirectly through some agonists.\(^3\) The contractile responses elicited by u46619 are also dependent significantly on activation of the L-type Ca\(^{2+}\) channel. Inhibition of the L-type Ca\(^{2+}\) channel led to a 50% reduction of maximal force production generated by u46619 in porcine coronary arteries.\(^3\) In the present study, expression of the L-type Ca\(^{2+}\) channel was down-regulated at both day points when compared with control and mock-infected vessels (Figure 3, A and B). This is similar to the contractile changes exhibited in response to KCl and u46619. IP3R, found on the sarcoplasmic reticulum (SR), is also involved in MLCK activation through the release of SR Ca\(^{2+}\) stores in response to agonist-mediated G-protein–coupled signaling.\(^3\) IP3R expression in infected coronary vessels remained unaffected at day 5 postinfection but was significantly down-regulated at day 10 postinfection (Figure 3, C and D). Because a decrease in contraction at day 5 postinfection was observed without a change in IP3R expression, and because a decrease in IP3R expression at day 10 postinfection did not result in a further reduction in contractile performance of infected tissue, the contribution of IP3R in the defects exhibited in response to u46619 is questionable. The most likely mechanism for the alterations in contractile responses to both high K\(^+\) and u46619 is through the reduction in L-type Ca\(^{2+}\) channel expression induced by C. pneumoniae infection. A decrease in the L-type Ca channel expression occurring in synthetic VSMCs is accompanied with an attenuation in smooth muscle actin and smooth muscle myosin heavy chain expression, two key components of the contractile apparatus. In the current study, although not significant, a decreasing trend in smooth muscle actin and smooth muscle myosin heavy chain expression in C. pneumoniae–infected vessels can be observed. Considering the mixed population of VSMCs (contractile and synthetic), it is possible the

![Figure 7. Superoxide detection by dihydroethidium staining in infected isolated coronary arteries. Representative staining of control, mock-infected, and Cpn-infected vessels was collected at day 10 postinfection. Fluorescence intensity (right panel) is represented as mean ± SEM, n = 4; *P < 0.05 versus control.](image-url)

![Figure 8. TLR-4, hHSP60, and cHSP60 localization within infected isolated coronary arteries. Representative staining of control (A–C), mock-infected (D–F), and Cpn-infected (G–I) vessels collected at day 10 postinfection. All panels: Hoescht, blue. A, D, G: TLR-4, red. B, E, H: hHSP60, green. C, F, I: cHSP60, green. Scale bar = 100 μm.](image-url)
abundant expression of these markers in the contractile VSMC population may be dampening the decreases in expression within the synthetic VSMC population, thus contributing to the overall expression observed within whole vessel lysates. As such, the potential contribution of both these contractile proteins in the altered contractile responses should not be completely dismissed.

Other mechanisms beyond simple changes in contractile machinery expression could be considered for the observed contractile responses. Structural changes in C. pneumoniae-infected vessels have previously been noted using this same model system. Synthetic VSMCs demonstrate an increase in extracellular matrix production. As such, an increase in medial thickening via VSMCs proliferation could, in theory, confer changes in vessel stiffness through increased collagen production. However, it is important to note that these potential alterations in vessel stiffness would likely occur at later time points as medial thickening only starts to occur from day 6 postinfection onward. It is, then, unlikely that the changes in vascular contractile function observed at day 5 postinfection are due to this phenomenon. In addition, the attenuation in contractile responses in C. pneumoniae-infected vessels is not further aggravated at day 10 postinfection, which suggests vessel stiffness is likely not a major contributor.

C. pneumoniae infection also induced a significant attenuation in bradykinin-induced relaxation of precontracted tissue (Figure 4B). Bradykinin elicits relaxation of vascular tissue through an endothelial-dependent mechanism. The impairment in vasorelaxation was limited to the endothelial layer in the present study, as C. pneumoniae infection did not affect relaxation induced by the nitric oxide (NO) donor sodium nitroprusside. C. pneumoniae is known to induce endothelial damage. However in this model, C. pneumoniae infection did not induce endothelial cell death. Therefore, the attenuated relaxation response is more likely due to a cellular mechanism and not a deterioration of the endothelial layer.

Alterations to the (eNOS)/NO pathway, leading to a reduction in NO bioavailability, represent the most important mechanism for impaired endothelial-dependent relaxation. Dihydroethidium staining of coronary vessels revealed an increase in superoxide production in both mock-infected and C. pneumoniae-infected vessels (Figure 7). Among the many sources of reactive oxygen species, NADPH oxidase is a major producer of superoxide within the vessel wall. p22phox, a major component of NADPH oxidase, is expressed in ECs, VSMCs, adventitial fibroblasts and macrophages within human atherosclerotic plaques. Expression of p22phox expression was upregulated in both mock-infected and C. pneumoniae-infected vessels at day 10 postinfection (Figure 6B). Furthermore, its expression was identified in both endothelial and medial layers (Figure 6C). The ability for the heat-inactivated C. pneumoniae to also elicit this response suggests that this effect is lipopolysaccharide mediated. Wick et al described that in vivo lipopolysaccharide administration increases hHSP60 expression on the surface of vascular endothelial cells. Our results indicate that C. pneumoniae and mock infection result in an increase in hHSP60 expression along the endothelium. In addition to down-regulating eNOS, Chen et al also described the ability of CHSP60 to up-regulate oxidative stress machinery including p22phox. Because of a conserved sequence homology for HSP60 between species, it is possible that hHSP60 may also induce similar downstream cellular changes through TLR-4 signaling.

The maintenance of isolated coronary vessels in a tissue culture environment for up to 10 days does not significantly alter the functional integrity of these arteries. Thus, the alterations observed in the present study are directly dependent on exposure to C. pneumoniae and/or C. pneumoniae components. The findings of the current study clearly demonstrate that C. pneumoniae infection is able to directly induce a significant impairment to the functional capacity of arteries. These defects are observed before (at 5 days) and after (10 days) significant thickening (hypertrophy) of the aorta is observed. These changes may have physiological consequences with regard to regulating normal blood flow, and may play an important role in...
the pathophysiology of hypertension, restenosis, and atherosclerosis.

References