Activation of the Protein Kinase C–Mediated Contractile System in Canine Basilar Artery Undergoing Chronic Vasospasm

Tohru Matsui, MD; Makoto Sugawa, MSc; Hiroo Johshita, MD; Yoh Takuwa, MD; and Takao Asano, MD

We previously suggested that activation of the protein kinase C–mediated contractile system may participate in the occurrence of chronic cerebral vasospasm. In the present study, we compared segments of normal beagle basilar arteries in vitro with segments of arteries undergoing chronic vasospasm to determine the responsiveness to various agonists such as serotonin, prostaglandin E₂, and phorbol 12,13-diacetate as well as to external Ca²⁺. We also compared the effects of W-7 (a calmodulin inhibitor), nicardipine (a calcium channel blocker), and H-7 (a protein kinase C inhibitor) on the spontaneous tonus of arterial segments stabilized at a resting tension of 3 g. Compared with normal segments, the responsiveness to each agonist in segments undergoing vasospasm was essentially unchanged whereas the responsiveness to external Ca²⁺ was significantly decreased (p<0.001). In segments undergoing vasospasm the decrease in resting tension induced by W-7 was markedly diminished (p<0.01), that induced by nicardipine was unchanged, and that induced by H-7 was significantly increased (p<0.01). Our results indicate that spontaneous tonus due to activation of the protein kinase C system is significantly augmented in segments undergoing vasospasm. Thus this system, rather than the Ca²⁺/calmodulin system, appears to play a major role in the occurrence of chronic vasospasm.

tively. Eight of 12 dogs in the first and fourth experiments and four of eight dogs in the second experiment were subjected to two-hemorrhage SAH.

The SAH was induced as previously reported. Under barbiturate anesthesia, cerebral angiography via the vertebral artery was carried out just prior to the induction of SAH on day 0 (control). On day 7, angiography was repeated and the dog was killed by exsanguination. The basilar artery was excised and immediately immersed in Krebs-Henseleit solution at room temperature. Arteries obtained from normal beagles and those exposed to two-hemorrhage SAH were dissected under an operating microscope and cut into six ring segments 5 mm long. Each segment was mounted on rigid prongs inside a chamber, which was then filled with 5 ml of Krebs-Henseleit solution (NaCl 120, KCl 4.5, CaCl₂ 2.5, MgCl₂ 1.0, NaHCO₃ 27, KH₂PO₄ 1.0, and glucose 10 in millimolar concentrations; pH 7.4±0.05). The Krebs-Henseleit solution was aerated with a 5% CO₂/95% O₂ gas mixture and kept at 37°C. The segments were allowed to equilibrate for 120 minutes before the start of each experiment while increasing stretch was applied to stabilize at a basal tension of 3 g. Changes in isometric tension following drug application were monitored by transducers (Nihon Kohden TB611, Japan) connected to a six-channel polygraph recorder (Nihon Kohden MC 6000).

In the first experiment, responses to serotonin, prostaglandin F₂α (PGF₂α), and PDA were compared between arterial segments from normal beagles (normal segments) and arterial segments from beagles exposed to two-hemorrhage SAH (spasm segments). Each test agent was added to the chamber in a cumulative manner. Changes in isometric tension following drug application were monitored by transducers (Nihon Kohden TB611, Japan) connected to a six-channel polygraph recorder (Nihon Kohden MC 6000).

In the third experiment, normal segments were stretched to stabilize at resting tensions of 1, 2, 3, or 4 g in chambers containing aerated Krebs-Henseleit solution. At each resting tension 10⁻⁶ M nicardipine, 10⁻⁵ M H-7, and 10⁻⁴ M papaverine were added to the chamber in a cumulative fashion, and the subsequent changes in resting tension were recorded.

In the fourth experiment, normal and spasm segments were stretched and stabilized at a resting tension of 3 g in aerated Krebs-Henseleit solution. The decrease in tension following the cumulative additions of 10⁻³ M W-7, 10⁻⁶ M H-7, and 10⁻⁴ M papaverine was recorded by the chamber in a cumulative fashion and the subsequent changes in resting tension were recorded.

In the first and fourth experiments, the spasm segments were further divided into moderate spasm segments (basilar artery diameter >60% of control) and severe spasm segments (basilar artery diameter <60% of control) according to the results of angiography carried out on day 7, prior to arterial excision. In the second experiment, only normal and severe spasm segments were used.

Values are expressed as mean±standard deviation. Dunnett's method or the Mann-Whitney test was used for statistical evaluation. Probability values less than 0.05 were considered to be significant.

Serotonin, diphenhydramine hydrochloride, atropine sulfate, R24571, and PGF₂α were pur-

<table>
<thead>
<tr>
<th>Table 1. Maximum Tension Developed in Response to Various Agents by Canine Basilar Artery Segments</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Segments</strong></td>
</tr>
<tr>
<td><strong>Agent</strong></td>
</tr>
<tr>
<td>Serotonin</td>
</tr>
<tr>
<td>Prostaglandin F₂α</td>
</tr>
<tr>
<td>Phorbol 12,13-diacetate</td>
</tr>
<tr>
<td>Ca²⁺⁺</td>
</tr>
<tr>
<td>Ca²⁺⁻</td>
</tr>
</tbody>
</table>

Ca²⁺⁺, Krebs-Henseleit solution; Ca²⁺⁻, calcium-free Krebs-Henseleit solution. *p<0.01 different from normal by Dunnett's method.
TABLE 2.  ED₉₀ Value for Various Agents in Canine Basilar Artery Segments

<table>
<thead>
<tr>
<th>Agent</th>
<th>Normal</th>
<th>Moderate spasm</th>
<th>Severe spasm</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>ED₉₀</td>
<td>n</td>
</tr>
<tr>
<td>Serotonin</td>
<td>18</td>
<td>2.73±1.5x10⁻⁴</td>
<td>22</td>
</tr>
<tr>
<td>Prostaglandin₁₂₋₁₃-diacetate</td>
<td>19</td>
<td>4.05±2.34x10⁻⁷</td>
<td>16</td>
</tr>
<tr>
<td>Ca(+)</td>
<td>10</td>
<td>9.5±4.21x10⁻⁸</td>
<td>22</td>
</tr>
<tr>
<td>Ca(−)</td>
<td>10</td>
<td>2.19±0.75x10⁻⁸</td>
<td>22</td>
</tr>
<tr>
<td>CaCl₂</td>
<td>12</td>
<td>2.5±1.12x10⁻⁴</td>
<td>18</td>
</tr>
</tbody>
</table>

Ca(+), Krebs-Henseleit solution; Ca(−), calcium-free Krebs-Henseleit solution.

*p<0.001 different from normal by Dunnett’s method.

Results

In the first experiment, diameters of the moderate and severe spasm segments were 68.9±7.8% and 46.3±4.8% of control, respectively. As shown in Table 1, the maximum tension (Tmax) developed in response to serotonin or PDA was significantly less in the severe spasm segments than in the normal segments (p<0.01). The Tmax developed in response to PGF₂α or CaCl₂ was slightly reduced in the severe spasm segments, although the difference was not significant. The ED₉₀ values for each agent are shown in Table 2. Although the ED₉₀ values for serotonin, PGF₂α, and PDA did not change significantly, that for CaCl₂ increased significantly in the moderate and severe spasm segments (p<0.001), with a shift of the dose–response curve to the right.

Among the various agents tested, only the protein kinase C inhibitor H-7 had a significant and dose-dependent inhibitory effect on the contraction elicited by PDA (Figure 1). The normal and spasm segments showed similar responses to H-7. None of the neurotransmitter antagonists or calmodulin inhibitors (R24571 and W-7) or the calcium channel blocker (nicardipine) significantly inhibited PDA-induced contraction.

Figure 2 shows the decreases in isometric tension following the cumulative applications of nicardipine and H-7 at different initial resting tensions. The final addition of papaverine induced no further decrease. The decrease in tension induced by nicardipine tended to increase as the initial tension increased, becoming maximal at a beginning tension of 3 g. H-7 also caused a significant decrease in tension, although the decrease was considerably less than that induced by nicardipine. An abrupt increase in the decrease in tension induced by H-7 was observed at a resting tension of 3 g.

Table 3 shows the decrease in tension following the cumulative applications of W-7, nicardipine, and H-7. Compared with the normal segments, severe spasm segments showed a marked diminution in the response to W-7 (p<0.01). The response to nicardipine was similar among the groups. A highly significant (p<0.01) increase in the decrease in tension induced by H-7 was revealed in both the moderate and severe spasm segments. No significant difference in the total reduction of tension after the addition of all three agents was revealed among the groups.

![Figure 1. Effects of various agents on contraction induced by phorbol 12,13-diacetate (PDA) in normal (top) and severe spasm (bottom) segments of canine basilar artery. 1, 10⁻⁶ M methysergide; 2, 10⁻⁸ M atropine; 3, 3x10⁻⁶ M diphenhydramine; 4, 10⁻⁴ M phentolamine; 5, 10⁻⁶ M R24571; 6, 10⁻³ M W-7; 7, 10⁻⁸ M nicardipine; 8, 10⁻³ M H-7; 9, 10⁻⁴ M H-7. Representative tracings from normal and spasm segments. Note decrease in tension to below baseline after addition of 10⁻⁴ M H-7.](http://stroke.ahajournals.org/)

![Figure 2.](http://stroke.ahajournals.org/)
Figure 2. Drop in tension induced by cumulative additions of nicardipine and H-7 at each initial resting tension in canine basilar artery segments. Data for papaverine are not shown because that agent induced no further drop in tension.

Discussion

In agreement with preceding reports, our study showed that the $T_{\text{max}}$ values for serotonin, PGF$_{2\alpha}$, and PDA were reduced in severe spasm segments whereas the respective $E_{\text{D}50}$ values remained unchanged. Hence, the responsiveness to each agent appears to remain essentially intact even in segments with severe vasospasm. It is noteworthy that spasm segments showed a diminished responsiveness to external Ca$^{2+}$ ($p<0.001$). Later discussion will be directed to the possible underlying mechanism.

In a previous study, in both normal and spastic segments contraction induced by PDA was inhibited by H-7 but not by the remaining agents, such as various antagonists to neurotransmitters, W-7, R24571, and nicardipine. That in vivo study, using the same battery of agents that applied directly to canine basilar artery segments undergoing chronic vasospasm, provided results similar to ours. Such an analogy between the results of in vitro and in vivo experiments lends support to the view that chronic vasospasm represents sustained smooth muscle contraction due to activation of protein kinase C.

In our fourth experiment, arterial segments were stabilized at a resting tension of 3 g, which is ascertained to give optimal dilatory responses to pharmacological agents. The response to W-7 was markedly diminished in severe spasm segments ($p<0.01$), underscoring the report of Sakaki et al. These data could be taken as an indication that the Ca$^{2+}$/calmodulin system does not play a pivotal role. In this regard, it is intriguing to note that the spasm segments showed a highly significant ($p<0.01$) increase in the decrease in tension induced by H-7. These data clearly indicate that spontaneous tonus due to protein kinase C activation is augmented in segments undergoing chronic vasospasm.

Protein kinase C activation within the canine basilar artery undergoing vasospasm is also supported by biochemical data showing a significant increase in the arterial content of 1,2-diacylglycerol, an intrinsic activator of the enzyme. Recently, we have shown that membrane translocation of protein kinase C is also significantly increased in the canine basilar artery (unpublished data). Thus, the results of our present study may be interpreted as indicating that activation of the protein kinase C system leads to augmentation of spontaneous tonus within the cerebral arteries.

Based on the above findings, the diminished contractile response to external Ca$^{2+}$ in the spasm segments deserves an attempt at explanation. First, the diminished availability of calmodulin hinders myosin light chain kinase activation even if the function of

### Table 3. Reduction in Resting Tension Induced by Various Agents in Canine Basilar Artery Segments

<table>
<thead>
<tr>
<th>Agent</th>
<th>Normal Resting Tension</th>
<th>Moderate spasm Resting Tension</th>
<th>Severe spasm Resting Tension</th>
</tr>
</thead>
<tbody>
<tr>
<td>W-7</td>
<td>0.384±0.135</td>
<td>0.486±0.152</td>
<td>0.002±0.095*†</td>
</tr>
<tr>
<td>Nicardipine</td>
<td>0.399±0.145</td>
<td>0.524±0.191</td>
<td>0.310±0.135</td>
</tr>
<tr>
<td>H-7</td>
<td>0.173±0.077</td>
<td>0.298±0.076*</td>
<td>0.300±0.094*</td>
</tr>
<tr>
<td>10°4 M</td>
<td>0.007±0.023</td>
<td>0.070±0.026†</td>
<td>0.111±0.078*</td>
</tr>
<tr>
<td>10°5 M</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>0.965±0.825</td>
<td>1.378±0.378</td>
<td>0.723±0.471</td>
</tr>
</tbody>
</table>

Values are mean±SD g.

* $p<0.01$ and 0.05, respectively, different from normal by Dunnett's method.
† $p<0.01$ different from moderate spasm by Dunnett's method.
the calcium channel is unaffected. Second, activation of protein kinase C, leading to stimulation of plasma membrane Ca\(^{2+}\)-ATPase and Na\(^+\)/Ca\(^{2+}\) exchange, causes suppression of the calcium-signaling pathway through an increased extrusion of intracellular Ca\(^{2+}\).\(^{18}\) Whereas these events are considered to be causally related to the diminished responsiveness of spasm occurrence of chronic vasospasm.\(^{19,20}\) Further investigation may be required regarding the above issue.

In summary, our present study shows that compared with normal segments, the responsiveness of spasm segments to various agonists such as serotonin, PGF\(_{2\alpha}\) and FDA was essentially unchanged whereas that to external Ca\(^{2+}\) was decreased. In the spasm segments, on the other hand, the portion of spontaneous tonus attributable to protein kinase C activation was significantly augmented. The results of this study, together with the already reported in vivo data,\(^{12}\) indicate that activation of the protein kinase C system, rather than of the Ca\(^{2+}\)/calmodulin system, plays a major role in the occurrence of chronic vasospasm.

**Acknowledgments**

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**References**

7. Varsos VG, Lisetzak TM, Han DH, Kistler JP, Vielma J, Black PM, Heros RC, Zervas NT: Delayed cerebral vasospasm is not reversed by aminophylline, nifedipine, or papaverine in a 'two-hemorrhage' canine model. *J Neurosurg* 1988;71:11-17

**Key Words** • cerebral vasospasm • protein kinase C • subarachnoid hemorrhage • dogs
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