Prolyl Oligopeptidase Is Involved in Release of the Antifibrotic Peptide Ac-SDKP

Maria A. Cavasin, Nour-Eddine Rhaleb, Xiao-Ping Yang, Oscar A. Carretero

Abstract—N-acetyl-seryl-asparyl-lysyl-proline (Ac-SDKP) is a ubiquitous tetrapeptide hydrolyzed almost exclusively by angiotensin-converting enzyme (ACE). Chronic treatment with Ac-SDKP decreases cardiac and renal fibrosis and inflammatory cell infiltration in hypertensive rats. However, very little is known about endogenous synthesis of Ac-SDKP, except that thymosin-β₄ may be the most likely precursor. Two enzymes are potentially able to release Ac-SDKP from thymosin-β₄: prolyl oligopeptidase (POP) and endoproteinase asp-N. POP is widely present and active in several tissues and biological fluids, whereas endoproteinase asp-N appears to be lacking in mammals. Therefore, we hypothesized that POP is the main enzyme involved in synthesizing the antifibrotic peptide Ac-SDKP. We investigated in vitro and in vivo production of Ac-SDKP. Using kidney cortex homogenates, we observed that Ac-SDKP was generated in a time-dependent manner in the presence of exogenous thymosin-β₄, and this generation was significantly inhibited by several POP inhibitors (POPi), Z-prolyl-proline, Fmoc-prolyl-pyrrolidine-2-nitrile, and S17092. Long-term administration of S17092 in rats significantly decreased endogenous levels of Ac-SDKP in the plasma (from 1.76±0.2 to 1.01±0.1 nM), heart (from 2.31±0.21 to 0.83±0.09 pmol/mg protein), and kidneys (from 5.62±0.34 to 2.86±0.76 pmol/mg protein). As expected, ACE inhibitors significantly increased endogenous levels of Ac-SDKP in the plasma, heart, and kidney, whereas coadministration of POPi prevented this increase. We concluded that POP is the main enzyme responsible for synthesis of the antifibrotic peptide Ac-SDKP. (Hypertension. 2004;43:1140-1145.)

Key Words: angiotensin-converting enzyme ■ heart ■ kidney

Ac-SDKP is a ubiquitous tetrapeptide normally present in organs and biological fluids of humans and experimental animals.¹⁻³ It was originally described as a natural inhibitor of pluriplotent hematopoietic stem cells, preventing entry into the S phase.⁴ Recently, Ac-SDKP was found to be hydroyzed almost exclusively by angiotensin-converting enzyme (ACE). After treatment with ACE inhibitors (ACEi), its plasma concentration increased 5-fold.⁵⁻⁷ Thus, some of the protective effects of ACEi may be mediated by an increase in Ac-SDKP. Our laboratory has recently reported that Ac-SDKP prevents fibroblast proliferation and collagen synthesis in vitro⁸ and that long-term treatment decreases cardiac and renal fibrosis and inflammatory cell infiltration in hypertensive rats without altering blood pressure.⁹,¹⁰ However, very little is known about endogenous synthesis of Ac-SDKP and the enzymes involved in the release of this tetrapeptide from its precursor thymosin-β₄. Thymosin-β₄, the most abundant β-thymosin in mammals,¹¹ was reported to be the most likely precursor of the tetrapeptide because it possesses the sequence Ac-SDKP in its N-terminus, and thus a single cleavage (Pro⁴-Asp⁵) would be sufficient to release the tetrapeptide.¹²,¹³ However, we could find no reports addressing the enzymes involved in the production of Ac-SDKP in vivo. Two enzymes are potentially able to cleave the Pro⁴-Asp⁵ bond of thymosin-β₄ to release Ac-SDKP: prolyl oligopeptidase (POP), which cleaves the carboxyl end of proline, and endoproteinase asp-N, which cleaves the amino end of aspartate. Lenfant et al previously reported that endoproteinase asp-N is the main enzyme responsible for Ac-SDKP synthesis, based on in vitro studies with a bacterial enzyme.¹² However, we know of no evidence that endoproteinase asp-N is present in mammals, which would suggest that it plays little or no role in production of Ac-SDKP in vivo. The other candidate enzyme, POP, belongs to a new class of serine peptidases (clan SC, family S9) unrelated to the well-known trypsin and subtilisin families¹⁴ and is able to hydrolyze a wide variety of proline-containing peptides.¹⁵ Unlike endoproteinase asp-N, POP is widely present and active in human and animal tissues and biological fluids.¹⁶⁻¹⁹ Therefore, we hypothesized that POP is the main enzyme responsible for release of the antifibrotic peptide Ac-SDKP. We tested whether POP is involved in release of Ac-SDKP from thymosin-β₄ in vitro, and whether treatment with a specific POP inhibitor (POPi) would lower endogenous levels of Ac-SDKP in vivo.

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Methods

Animals
We used 10-week-old male Sprague-Dawley rats (Charles River Laboratories). All surgical procedures were conducted with the animals under pentobarbital anesthesia (50 mg/kg IP). The study was approved by the Institutional Animal Care and Use Committee (IACUC) of Henry Ford Health System.

Ac-SDKP Determinations
Ac-SDKP in plasma, urine, and tissue was measured as described by Azizi et al.16 using a commercially available enzyme immunoassay kit (Cayman Chemicals). Blood was collected in a prechilled syringe containing 10 µL heparin and 10 µL 10^{-3} mol/L lisinopril per mL of blood, and the plasma was separated. The purpose of adding an ACEi was to prevent Ac-SDKP degradation during sample processing. Urine was collected directly from the bladder in a prechilled syringe containing 10 µL 10^{-3} mol/L lisinopril. Tissue samples were homogenized with cold 50 mmol/L phosphate-saline buffer (pH=7.4) containing 10^{-3} mol/L lisinopril. Ac-SDKP was extracted from an aliquot of plasma or tissue homogenate (150 and 100 µL, respectively) and processed according to the manufacturer’s instructions. Urine was assayed without precipitation in 1:100 dilutions. We observed no cross-reactivity with thymosin-β_4. Total proteins were measured by the Bradford method and creatinine by the Jaffé reaction.

Effect of POPi on Ac-SDKP Synthesis In Vitro
Homogenates of kidney cortex were chosen because we have evidence that Ac-SDKP has renal antifibrotic effects.10 Tissue was desegregated and diluted up to 150 µg/mL total protein with cold phosphate–saline buffer; 100-µL aliquots of the homogenate were incubated at 37°C with: (1) thymosin-β_4 vehicle and POPi vehicle (time control); (2) thymosin-β_4 (2 µmol/L) plus POPi vehicle; (3) thymosin-β_4 plus POPi (0.01 µmol/L); or (4) thymosin-β_4 plus POPi (0.1 µmol/L). Thymosin-β_4 purified from bovine spleen (gift of D. Safer, University of Pennsylvania, Philadelphia, PA),20 was dissolved in water. Three different POPi were used: Z-prolyl-prolinal (Z-Pro-Pri), Fmoc-prolyl-pyrrolidine-2-nitrile (Fmoc-Pro-PyrrCN), and thymosin-β_4. Data are expressed as mean±SE.

Results

Effect of POPi on Ac-SDKP Synthesis In Vitro
Figure 1 shows time-dependent production of Ac-SDKP in vitro. Kidney cortex homogenates were incubated with exogenous thymosin-β_4 in the presence or absence of different POPi concentrations. We used a higher dose of S17092 than in the previous protocol to ensure maximum effect. After 5 days of treatment, rats were anesthetized and the femoral artery and vein catheterized for mean arterial pressure (MAP) measurement and drug administration, respectively. After a 20-minute stabilization period, MAP was recorded for 5 minutes (basal), followed by different doses of bradykinin (Bk) (25, 50, and 100 ng) or angiotensin I (Ang I) (12, 25, and 50 ng). After each dose, MAP was allowed to stabilize for 15 minutes.

Data Analysis
In vitro experiments and changes in MAP were analyzed using ANOVA, and the Fisher protected least significant difference (Fisher LSD) was used to adjust for multiple comparisons. Ac-SDKP levels in vivo were analyzed using the Wilcoxon nonparametric rank-sum test. P<0.05 was considered significant. Data are expressed as mean±SE.
POPi (n=7). Panels A, B, and C show production of Ac-SDKP in the presence of Z-Pro-Pro, Fmoc-Pro-Pyr-CN, and S17092, respectively. When homogenates were incubated in the presence of exogenous thymosin-β4, Ac-SDKP increased significantly in a time-dependent manner compared with time control (all panels). The highest concentration of Z-Pro-Pro (0.1 mol/L) significantly blocked Ac-SDKP synthesis, whereas a lower dose (0.01 mol/L) was ineffective (panel A); 0.1 mol/L Fmoc-Pro-Pyr-CN significantly inhibited Ac-SDKP production, whereas 0.01 mol/L gave only partial inhibition (panel B). S17092 at high and low concentrations completely inhibited Ac-SDKP synthesis (panel C).

Effect of an Oral POPi on Endogenous Ac-SDKP

Figure 2 shows concentration of Ac-SDKP in plasma and urine, and Figure 3 shows Ac-SDKP content in the heart and kidney after 7 days of S17092 treatment. Plasma levels of Ac-SDKP were slightly lower in rats treated with S17092 alone compared with vehicle but did not reach statistical significance, whereas urinary levels were similar in both groups. Plasma and urinary Ac-SDKP increased significantly after captopril treatment, and this increase was significantly prevented by coadministration of S17092. Chronic treatment with S17092 alone significantly decreased endogenous levels of Ac-SDKP in the heart and renal cortex compared with vehicle. Captopril significantly increased endogenous Ac-SDKP, whereas coadministration of S17092 significantly prevented this increase.

Effect of S17092 on ACE Activity In Vivo

We tested whether a high dose of S17092 can inhibit ACE. In this protocol, we doubled the dose of S17092 used to measure endogenous Ac-SDKP to maximize possible effects of POPi on ACE activity and avoid problems related to insufficient dosage. Figure 4 shows changes in MAP caused by bolus injection of increasing amounts of Bk (panel A) and Ang I (panel B) in rats treated with vehicle, captopril, or S17092. Because Bk and Ang I are ACE substrates, they were used as indicators of ACE activity. We found that in rats treated with captopril, Bk-induced hypotension was significantly enhanced and the Ang I-induced pressor effect was significantly inhibited compared with vehicle. However, the changes in MAP seen in rats treated with S17092 were no different from vehicle. Plasma Ac-SDKP was also measured after the experiments, and we found that POPi alone at 40 mg/kg per day significantly decreased Ac-SDKP compared with vehicle (1.76±0.2 vs. 1.01±0.1 nM; P<0.05).

Discussion

We found that: (1) Ac-SDKP generation in kidney homogenates increased in a time-dependent manner in the presence of exogenous thymosin-β4, and this process was blocked significantly by several POPi; (2) rats that received POPi long-term showed significantly lower levels of endogenous Ac-SDKP in the heart and kidney; and (3) ACEi significantly increased endogenous Ac-SDKP in the plasma, urine, heart, and kidney, whereas coadministration of POPi significantly prevented this increase. We believe this is the first study showing that POP is the enzyme responsible for production of the antifibrotic peptide Ac-SDKP in vitro and in vivo. Thymosin-β4 was postulated to be the most likely precursor for the tetrapeptide, because it possesses the sequence Ac-.
SDKP in its N-terminus. Thymosin-β₄ was originally found in the thymus but is widely present in mammals and distributed ubiquitously in tissue and in the circulation. Its structure is the same in all mammals studied so far, including humans, rats, dogs, mice, cats, calves, pigs, sheep, and horses, except for the rabbit, which has Ala⁴ instead of Pro⁴. Although we have no direct evidence that Ac-SDKP is released from thymosin-β₄ in vivo, precursor and product peptides have been colocalized to many tissues and their concentration shows a positive correlation. Moreover, in the present study we found that addition of exogenous thymosin-β₄ increased Ac-SDKP production in vitro, supporting this assumption.

Lenfant et al previously reported that endoproteinase asp-N was the enzyme responsible for release of Ac-SDKP from thymosin-β₄ in vitro.¹² They arrived at this conclusion by incubating pure bacterial endoproteinase asp-N and POP with 1 mg thymosin-β₄ for 18 hours. However, we believe the in vitro conditions used in their study are questionable for the following reasons: (1) bacterial enzymes could not properly reproduce the in vivo mammalian enzymatic and biochemical environment; and (2) incubation of extremely high amounts of substrate for a long time could lead to loss of enzyme specificity. In the present study, we clearly demonstrated that POP is responsible for Ac-SDKP production both in vitro and in vivo, because generation of the tetrapeptide was significantly blocked and endogenous levels were significantly decreased by specific POP inhibitors.

One particular characteristic of POP is that it only cleaves oligopeptides no larger than 30 amino acids. Its catalysis is controlled by a gating filter mechanism that only allows small peptides to gain access to the active site; in this way, larger structural peptides and proteins are protected from proteolysis. Because thymosin-β₄ has 43 amino acids, we believe it must be hydrolyzed into a smaller peptide by some other peptidases, resulting in truncation of thymosin-β₄ before POP hydrolysis and release of Ac-SDKP from the N-terminus. Although we have not tested whether the real substrate for POP would be a truncated rather than an intact thymosin-β₄, it is still important to show that POP is responsible for biosynthesis of Ac-SDKP. This could explain why in Lenfant’s studies of pure POP did not generate Ac-SDKP from thymosin-β₄, because the incubates only contained the pure bacterial POP and thymosin-β₄, and the absence of other peptidases made it impossible to produce a truncated thymosin-β₄ as a substrate for POP and finally release Ac-SDKP. In contrast, our in vitro studies using tissue homogenates have the advantage of providing the proper enzymatic environment for hydrolysis and release of Ac-SDKP from thymosin-β₄, because peptidases normally found in the kidney are still present in the homogenate.

We blocked Ac-SDKP synthesis using 3 different POPi, which have been developed to treat mental disorders such as depression, anorexia and bulimia nervosa, and Alzheimer disease. In general, peptide aldehyde analogs of the acyl portion of protease substrates are potent competitive inhibitors of serine and sulfhydryl proteases, because the aldehyde interacts with the active site serine to form a tetrahedral transition-state analog. Based on this, S. Wilk and M. Orlowski synthesized Z-prolyl-prolinal (Ki 5 nM).²⁸ Fmoc-Pro-PyrCN (Kᵢ 5 nM) was also synthesized by S. Wilk;²¹ contrary to Z-pro-prolinal, this is a reversible inhibitor that could easily be synthesized from commonly available intermediates using a simple 2-step procedure. S17092 (Kᵢ 1.5 nM) was developed by the Institut de Recherches Internationale Servier²² and was described as a highly potent, specific, and cell-permeant inhibitor of proline endopeptidase. All of these inhibitors were tested for their capacity to block several other proteolytic enzymes at a concentration >1000-fold greater than their Kᵢ for POP, and all failed to do so. Although these inhibitors are reportedly highly specific and do not block other peptidases, by using several compounds we increased the degree of certainty that proteolytic release of Ac-SDKP from thymosin-β₄ is caused by POP.

Endogenous Ac-SDKP in the heart and kidney was significantly lower in rats treated with POPi alone (lower dose) compared with vehicle, but not in the plasma. However, plasma levels of Ac-SDKP were significantly lower when a higher dose was used. It could be argued that Ac-SDKP in plasma did not decrease significantly because plasma ACE activity is inhibited by POPi; if so, Ac-SDKP degradation is blocked and its concentration would not decrease (Figure 5 shows clarification of enzymatic reactions). Therefore, we tested whether ACE activity is affected during chronic POPi treatment. We found that in rats treated with ACEi, Bk-induced vasodilatation was enhanced and Ang I-induced vasoconstriction was inhibited; however, in rats treated with a high dose of S17092, changes in MAP caused by acute administration of Bk and Ang I were no different from vehicle, suggesting that ACE activity is not affected by POPi.
Generation and degradation of Ac-SDKP

**Thymosin β₄**: Ac-Ser-Asp-Lys-Pro-Asp ... (43 amino acids)

POPinh \(\rightarrow\) POP (prolyl oligopeptidase)

Ac-SDKP

ACEinh \(\rightarrow\) ACE

Inactive fragments

**Figure 5.** Schematic diagram illustrating enzymatic generation and degradation of Ac-SDKP.

This confirms the findings of Barelli et al, who tested a saturating concentration of S17092 in vitro and observed that peptidase activity, including ACE, was not affected.²²

Rats that received long-term ACEi treatment showed significantly increased endogenous levels of Ac-SDKP in the plasma, urine, heart, and kidney. This was as expected, because Ac-SDKP degradation is blocked by ACEi, and this was in accord with previous studies.⁵,⁶,²⁰ By increasing Ac-SDKP levels with an ACEi, we wanted to show that simultaneous inhibition of tetrapeptide synthesis by coadministration of POPi could prevent Ac-SDKP accumulation.

Peptidases that are able to hydrolyze adjacent proline bonds are considered to have great metabolic importance, because the presence of a proline residue confers certain conformational restrictions to the peptide chain along with increased resistance to hydrolysis.³⁰,³¹ POP is a member of the prolyl oligopeptidase family, a relatively new class of serine peptidases. It is involved in the maturation and degradation of many peptide hormones and neuropeptides such as angiotensins, kinins, vasopressin, and substance P,¹⁵,¹⁷ and is widely distributed in various tissues, ¹⁷,¹⁹ Ac-SDKP can be produced locally in any organ and released into the circulation. Ac-SDKP was also found to be synthesized by bone marrow cells in long-term murine³² and human ³³ cultures, although we know of no direct evidence showing POPi could prevent Ac-SDKP accumulation.

Because POP has been described as a ubiquitous enzyme that is widely distributed in various tissues,¹⁷,¹⁹ Ac-SDKP can be produced locally in any organ and released into the circulation. Ac-SDKP was also found to be synthesized by bone marrow cells in long-term murine ³² and human ³³ cultures, although we know of no direct evidence showing POP expression in bone marrow. We performed some preliminary experiments in vitro, incubating rabbit bone marrow homogenates with thymosin β₄, and found that production of Ac-SDKP was blocked in the presence of POP inhibitors, similar to what we report in the present article involving kidney homogenates. Although we decided to use the kidney in future experiments because of its relevance to our research, based on our preliminary data we believe POP is present in bone marrow.

**Perspectives**

Increased extracellular matrix deposition is an important feature of target organ damage in hypertension, and we have reported that Ac-SDKP reduces collagen deposition in the heart and kidney of hypertensive rats by inhibiting fibroblast proliferation and decreasing macrophage infiltration.⁹,¹⁰ However, biosynthesis of Ac-SDKP has been poorly studied. We found that POP is the major enzyme involved in endogenous production of Ac-SDKP. Therefore, future studies in vivo involving long-term inhibition of Ac-SDKP synthesis will help us determine: (1) whether Ac-SDKP plays a physiological role in antagonizing pro-fibrotic stimuli such as Ang II and endothelin-I; and (2) whether endogenous Ac-SDKP is an important mediator in the protective effects of ACEi in hypertension, acting as an anti-inflammatory cytokine to prevent fibrosis and inflammation. Our present findings will help us understand some of the mechanisms by which fibrosis develops in diseases characterized by collagen deposition, such as hypertension, diabetes, and heart failure.

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**References**


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