

Genomic Structure of the Amphioxus Calcium Vector Protein¹

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Calcium vector protein (CaVP) is an EF-hand Ca²⁺-binding protein, which is unique to the protochordate, amphioxus. CaVP is supposed to act as a Ca²⁺ signal transducer, but its exact function remains unknown. Not only its function but also its exact evolutionary relationship to other Ca²⁺-binding proteins is unclear. To investigate the evolution of CaVP, we have determined the complete sequences of CaVP cDNAs from two amphioxus species, *Branchiostoma lanceolatum* and *B. floridae*, whose open reading frame cDNA and amino acid sequences show 96.5 and 98.2% identity, respectively. We have also elucidated the structure of the gene of *B. floridae* CaVP, which is made up of seven exons and six introns. The positions of four of the six introns (introns 1, 2, 3, and 5) are identical with those of calmodulin, troponin C, and the Spec protein of the sea urchin. These latter proteins belong to the so-called troponin C superfamily (TnC superfamily) and thus CaVP likely also belongs to this family. Intron 6 is positioned in the 3' noncoding region and is unique to CaVP, so it may represent a landmark of the CaVP lineage only. The position of intron 4 is not conserved in the genes of the TnC superfamily or CaVP, and seems to result from either intron sliding or the addition of an intron (randomly inserted into or close to domain III) to the genes of the TnC superfamily during their evolution.

Key words: amphioxus, calcium vector protein, cDNA sequences, EF-hand.

The EF-hand is a Ca²⁺-binding motif normally composed of 29 amino acids with a helix-loop-helix structure (1). Present-day proteins possess two to eight EF-hands and are classified as belonging to one protein family, called collectively the EF-hand family. The family comprises 39 subfamilies (2), and four of them, calmodulin (CaM), troponin C (TnC), myosin essential light chain (ELC), and myosin regulatory light chain (RLC), called the CTER subfamily, are evolutionarily closely related according to the results of analyses at both the amino acid (3) and cDNA levels (4). In fact, such a close relationship has been suggested from the earliest reports (5, 6), and the CTER together with parvalbumin (7) and Spec (8-10), a Ca²⁺-binding protein isolated from the sea urchin, have often been called the "TnC superfamily". Except for parvalbumin, they possess four EF-hand domains per molecule and may have evolved from a common four-domain ancestor. It is thought that parvalbumin lost the first domain during its evolution.

Calcium vector protein (CaVP) is a unique protein found

in various tissues of amphioxus and is abundantly expressed in muscle (11). It forms a stable complex with a protein of 26 kDa, CaVP-target protein (CaVPT, 12-14), and it is supposed to act as a Ca²⁺ signal transducer, although its exact function remains unknown. The evolution of CaVP is also uncertain. Similar to the members of the TnC superfamily, CaVP possesses four EF-hand domains, but binds only two Ca²⁺ per molecule. EF-hands I and II do not bind Ca²⁺, and in the native protein, a unique disulfide bridge is formed between these two domains. Kobayashi *et al.* (11) stressed that CaVP shows about 35% sequence identity with CaM and TnC. At the same time, however, they also suggested the possibility that CaVP is a chimera with a C-terminal half homologous to that of CaM and TnC, and an N-terminal half resembling those of parvalbumins and sarcoplasmic Ca²⁺ binding proteins. On the other hand, CaVP clusters with ELC on the dendrogram constructed by the maximum parsimony method (15), and on analysis of the individual EF-hand domains, those of the CaVP were found to cluster near those of S100, the small EF-hand protein which possess two domains per molecule (3).

In this study, we determined the complete cDNA sequences of CaVPs from two species, *Branchiostoma lanceolatum* and *B. floridae*, and elucidated the exon/intron organization of the latter and discussed the evolutionary relationship of CaVP with the TnC superfamily.

MATERIALS AND METHODS

Cloning of B. lanceolatum CaVP cDNA—Total RNA of adult *B. lanceolatum* was prepared according to the acid guanidium thiocyanate method (16), and mRNA was

¹ This work was supported in part by a grant from the Japan Society for the Promotion of Science. The determined nucleotide sequences have been submitted to the DDBJ under the accession numbers, AB001688 (*Branchiostoma lanceolatum* CaVP cDNA), AB001689 (*B. floridae* CaVP cDNA), and AB023071-AB023073 (*B. floridae* CaVP genomic clones).

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Abbreviations: CaM, calmodulin; CaVP, calcium vector protein; ELC, myosin essential light chain; nt, nucleotide; RLC, myosin regulatory light chain; TnC, troponin C.

purified with Oligotex dT-30 Super (Japan Roche). The single-stranded cDNA was synthesized using a First-Strand cDNA Synthesis Kit (Pharmacia). The cDNA of *B. lanceolatum* CaVP was amplified by PCR using Ex Taq DNA polymerase (Takara). The redundant oligomer used for PCR was 5'-GA(AG)GA(AG)AA(AG)GA(CT)GA(AG)-TG(CT)ATGAA-3', which was designed based on the amino acid sequence, EEKDECMK (residues 12-19), of *B. lanceolatum* CaVP (11). The oligo-dT adaptor, 5'-GGGATCC-GAATTCT₁₇-3', was used as another primer. The 5' upstream of cDNA was determined as follows. The EcoRI-ended double-stranded cDNA was synthesized from mRNA using a TimeSaver cDNA Synthesis Kit (Pharmacia), and the EcoRI Cassette (Takara) was ligated at each end of cDNA. The 5' upstream region was amplified by PCR using the cassette-specific primer C1, 5'-GTACATATTGTCGT-TAGAACGCG-3', and 5'-CGAAGACAACACAGACTTT-ATTAC-3' (complementary to the sequence from nt 1057 to 1080).

Cloning of *B. floridae* CaVP cDNA—The λ ZAP II cDNA library of embryonic *B. floridae* was kindly provided by Dr. L.Z. Holland (Scripps Institute of Oceanography, University of California, San Diego). The *B. floridae* CaVP cDNA was also amplified by PCR using the cDNA library as a template. The primers used for 3'-half amplification were 5'-TGAGGGCGTTCAAGGTCTTC-3' (corresponding to the sequence from nt 275 to 294 of *B. lanceolatum* CaVP cDNA) and the T7 primer, 5'-TAATACGACTCACTATA-GGG-3', and those for 5'-half amplification were 5'-GAAT-CCTGTGGACTAACTTA-3' (complementary to the sequence from nt 487 to 506 of *B. lanceolatum* CaVP cDNA) and the T3 primer, 5'-ATTAACCTCACTAAAGGGA-3'.

Screening of a Genomic Library—The λ FIX II genomic library of *B. floridae* was also provided by Dr. L.Z. Holland, and was screened using the open reading frame of *B. lanceolatum* CaVP cDNA as a probe, because the identity between these two species, CaVP cDNAs is very high (96.5%, see below). The probe was labeled with DIG-DNA Labeling Mixture (Boehringer Mannheim) by PCR using the following forward primer, 5'-CTCTACCAGTCTTTC-AGTCC-3' (corresponding to the sequence from nt -29 to -10) and the reverse primer, 5'-GAATCCTGTGGACTA-ACTTA-3' (complementary to the sequence from nt 487 to 506). Hybridization and washing were carried out according to the manufacturer's instructions (Boehringer Mannheim), and positive clones were detected with a DIG Luminescent Detection Kit (Boehringer Mannheim).

DNA Sequencing—All PCR-amplified products and isolated genomic clones were subcloned to the pCR II plasmid vector (TA-cloning kit, Invitrogen) or pUC18 for sequencing. The nucleotide sequences were determined by the dideoxy chain termination method with a Dye Primer Cycle Sequencing Kit (Applied Biosystems) using an automated DNA sequencer (Applied Biosystems 373A).

RESULTS AND DISCUSSION

CaVP cDNAs from Two *Amphioxus* Species—The cDNA of *B. lanceolatum* CaVP was amplified by PCR and the complete sequence of 1,124 nucleotides was established from two overlapping fragments (Fig. 1). The open reading frame is composed of 489 nucleotides and encodes a protein of 162 amino acid residues including the initial Met.

Compared to the previously reported amino acid sequence of *B. lanceolatum* CaVP (11), the sequence of positions 152 to 155 is Lys-Lys-Ser-Lys instead of Lys-Ser-Lys-Lys. This region is Lys-rich and therefore the precise sequence is difficult to deduce from the results of peptide analysis. This and the fact that the same sequence is also observed in *B. floridae* CaVP (Fig. 1) led us to conclude that the previous report probably contained a peptide sequence error.

The cDNA of *B. floridae* CaVP is composed of 1,080 nucleotides and the open reading frame is 489 nucleotides long, encoding a protein of 162 amino acid residues including the initial Met, i.e. it is of the same length as in *B. lanceolatum* (Fig. 1). Within the coding regions, there are 17 nucleotide substitutions (96.5% identical) reflecting three amino acid differences (98.1% identical) between the CaVPs of the two species. Two of the three substitutions occur in domain II and the other at the C-terminus. Thus, there is likely no difference in Ca²⁺ binding between the two CaVPs, i.e. both bind two Ca²⁺ per molecule at EF-hand domains III and IV.

The identity between the two species' CaVPs is as high as those of CaMs, TnCs and skeletal actins, which are known to be highly conserved proteins (see Table I). However, mitochondrial proteins, such as cytochrome *b* and NADH dehydrogenase, of the two amphioxus species show equally high identity, as compared to the lower level of identity between the human and mouse mitochondrial proteins. Therefore, the high similarity of the two species' CaVPs may not reflect the conservatism of CaVP, but may rather be due to relatively recent divergence of *B. lanceolatum* and *B. floridae*.

Genomic Structure of the CaVP Gene—Using *B. lanceolatum* CaVP cDNA as a probe, we initially isolated three positive clones (clones 1, 2, and 3) from the genomic library of *B. floridae*. As clones 1 and 2 showed similar digestion patterns with several restriction enzymes, only clones 1 and 3 were used for further investigation. The nucleotide sequence of clone 1 was determined: it comprises 6,729 bp and spans the 5'-half of the CaVP gene up to Ile-90 (Fig. 2A). In this region, the CaVP gene consists of four exons divided by three introns (introns 1 to 3), and the nucleotide sequences of the exons exactly match those of the cDNA. All the introns start with GT and end with AG, and, according to the nomenclature³ of Kretsinger and Nakayama (17), the introns positions are -14/0, 1.01/1, and 2.13/1, respectively.

Clone 3 comprises 4,613 bp and spans the 3'-half of the CaVP gene starting from Ile-105 (Fig. 2A). In this region, the gene consists of three exons divided by two introns (introns 5 and 6); the position of the former intron is 4.21/1, and the last intron is inserted within the 3'-untranslated

³The positions of introns are indicated according to the nomenclature of Kretsinger and Nakayama (17). The first number corresponds to the EF-hand domain numbered sequentially from N to C. The second number (following the period) represents the residue number of the intron insertion in the EF-hand domain, which as a rule consists of 29 residues. The last number (following the slash) is the phase: 0 means that the intron lies between triplet codons, 1 between first and second nucleotides of the codon, and 2 between the second and third ones. For instance, 4.21/1 means insertion in domain IV, 21st residue and phase 1. -14/0 means phase 0, 14 residues before the beginning of domain I. 3+01/1 means phase 1, 1 residue beyond domain III, within the region between domain III and IV.

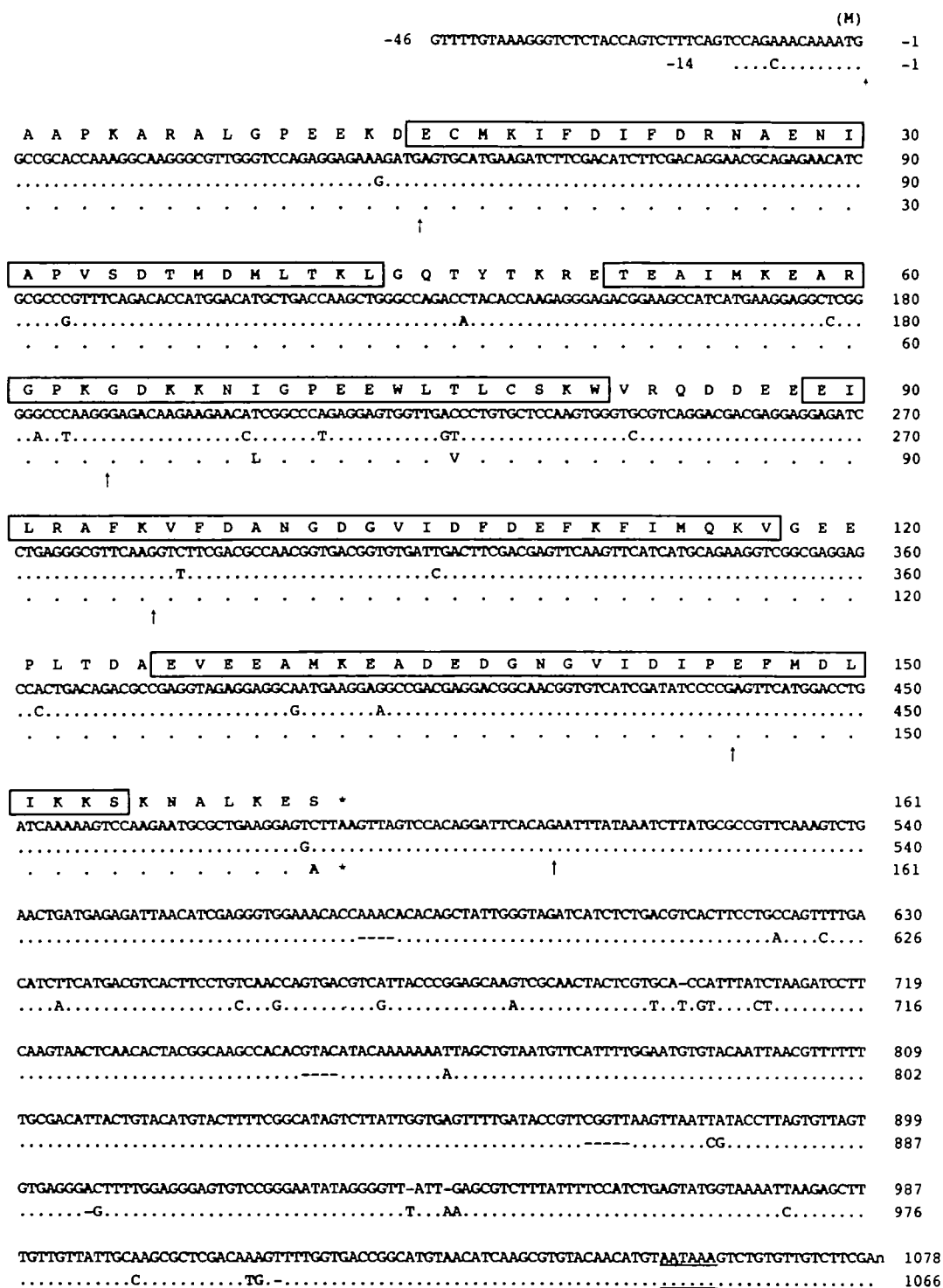


Fig. 1. Comparison of the cDNA and deduced amino acid sequences of CaVPs of two species of amphioxus. Upper lanes: *B. lanceolatum* CaVP cDNA and deduced amino acid sequence; lower lanes: *B. floridae* CaVP cDNA and deduced amino acid sequence. Identical nucleotides and amino acids to those in *B. lanceolatum* CaVP are indicated by dots (.), and gaps are inserted for maximal similarity

and shown by bars (-). The typical polyadenylation signals (AATAAA) are underlined, and the four EF-hand Ca²⁺ binding domains are boxed. The initiator methionine (residue -1, parenthesized) likely is removed after translation. The upward arrows (↑) indicate the positions of introns in the *B. floridae* CaVP genes.

region. Both introns are delineated in conformity to the GT/AG rule. Compared to the cDNA sequence, the genomic nucleotide sequence exhibits some differences: in the

open reading frame the GAT codon encoding Asp-143 changes to GAC in the genome and the 3'-noncoding region contains one substitution and three nucleotide insertions

(data not shown).

Since there is no overlapping region between clones 1 and 3, the region encoding from Leu-91 to Val-104 can not be adequately described. Therefore we performed further screening to obtain another clone containing the 91/104 region and succeeded in isolating clone 4. Clone 4 contains all the *CaVP* gene except exon 1 (see Fig. 2A). Of this clone only the exons and regions flanking the splice junctions were sequenced basically. Within the Leu-91/Val-104

region an intron (intron 4) was observed at position 3.08/0, whereas the positions of the other introns are identical to those in clones 1 and 3. The fourth intron was identified in clone 4 only, but in the *B. lanceolatum CaVP* gene an intron located at exactly the same position, as determined on sequencing of a partially amplified genomic PCR fragment (data not shown). To summarize, the conjectural organization of the *B. floridae CaVP* gene is shown in Fig. 2B, and it may consist of 7 exons and 6 introns (ca. 11.5 kbp).

When the intronic sequences of clones 1, 3, and 4 are compared, some substitutions and insertions/deletions are observed; the identity between each pair of splice junction flanking regions is ca. 90% on average. Moreover, in the exons four substitutions were observed, all of them being silent substitutions (data not shown). Thus, clone 4 may be derived from a gene that is different from in the case of clones 1 and 3. It is possible that clones 1 and 3 also come from distinct genes. There are two possible explanations for the observed discrepancies: (i) *amphioxus CaVP* may be multi-copy gene, or (ii) the different clones may be derived from distinct animal specimens, since the genomic library was not constructed from a single *amphioxus* specimen (18). Although the three clones are derived from different genes, it is supposed that the positions of the introns are identical in all genes.

Comparison of Intron Positions among TnC Superfamily Genes—The distribution of introns in the genes of *CaVP* and members of the TnC superfamily is shown in Table II. In all genes the first intron is basically inserted just after the initiation codon, ATG (shown as M in Table II). Furthermore, the positions of the second, third, and fifth

TABLE I. The identity (%) of several molecules between two species of *amphioxus*, and between human and mouse.

	<i>B. floridae</i> vs. <i>B. lanceolatum</i>		Human vs. Mouse	
	cDNA (ORF)	Amino acid	cDNA (ORF)	Amino acid
CaVP	96.5	98.2		
CaM	97.1	100.0	91.8	100.0
TnC	95.4	97.6	90.3	98.8
Skeletal actin	95.5	97.1	90.8	100.0
Cytochrome <i>b</i>	97.7	99.2	73.5	76.7
NADH 5 ^a	98.2	99.3	65.2	63.0

^aNADH 5, NADH dehydrogenase subunit 5. The accession numbers (EMBL/DBJ/GenBank) of the sequences used are; Y09863 (*B. floridae CaM*), Y09880 (*B. lanceolatum CaM*), M19311 (human *CaM II*), M27844 (mouse *CaM II*), D88977 (*B. floridae TnC*), D88976 (*B. lanceolatum TnC*), X07898 (human fast skeletal *TnC*), M57590 (mouse fast skeletal *TnC*), D87407 (*B. floridae* skeletal actin), Y13662 (*B. lanceolatum* skeletal actin), M20543 (human skeletal alpha-actin), M12347 (mouse skeletal alpha-actin), AF035173 (*B. floridae* cytochrome *b*), AF035172 (*B. floridae* NADH dehydrogenase subunit 5), Y16474 (*B. lanceolatum* mitochondrial DNA), D38112 (human mitochondrial DNA), and J01420 (mouse mitochondrial DNA).

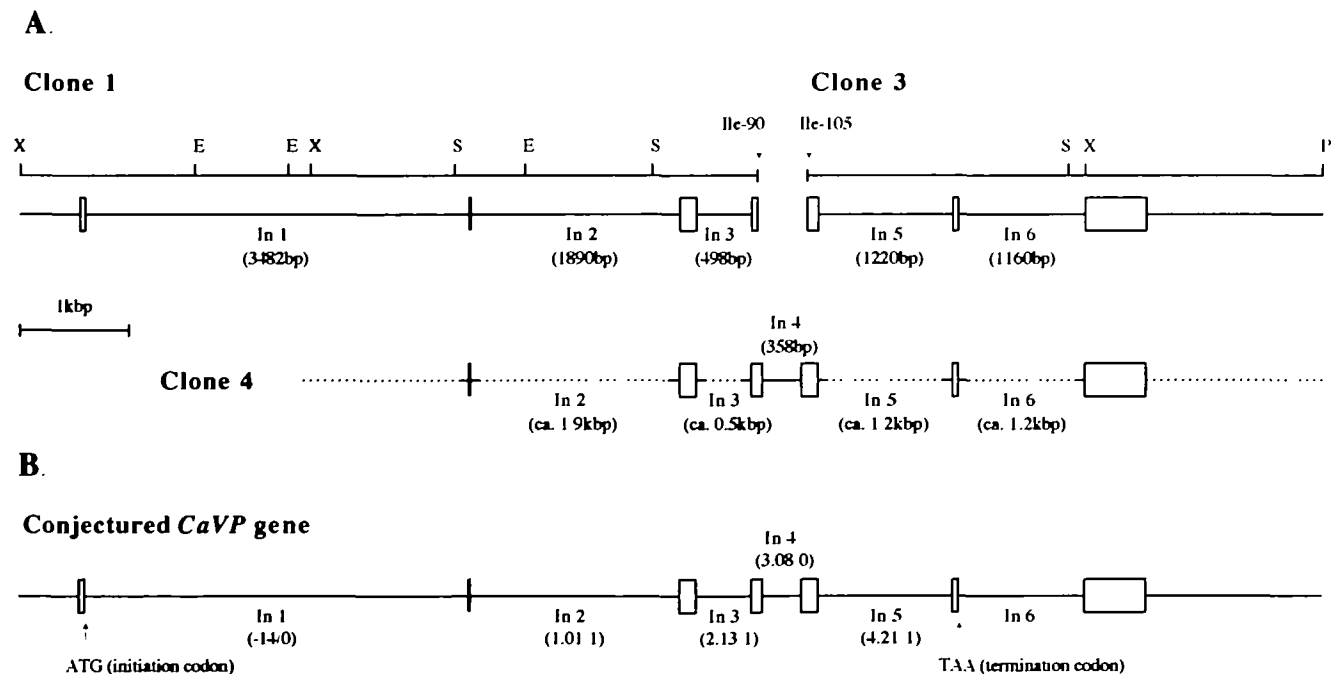


Fig. 2. Structural organization of the *B. floridae CaVP* genomic clones. A: Restriction and exon/intron maps of clones 1 and 3. *EcoRI* (E), *PstI* (P), *SaI* (S), and *XbaI* (X) restriction sites are indicated. Exons are shown by boxes, the 5'- and 3'-flanking regions and introns (In) are shown by bars, and the length of each intron is indicated. As for clone 4, the regions whose sequences were not

determined are shown by dotted lines, and the length of each intron was estimated by PCR. B: The conjectural organization of the gene of *B. floridae CaVP*. It may be composed of 7 exons and 6 introns, and the position of each intron is indicated in parenthesis. The positions of the initiation codon (ATG) and the termination codon (TAA) are also indicated.

TABLE II. The intron positions of TnC superfamily genes (based largely on Refs. 17 and 20).

Subfamily species (type)	Positions of introns					
CaVP						
Amphioxus	-14/0 (M)	1.01/1	2.13/1	3.08/0	4.21/1	3'-n.r.*
TnC						
Human (f)	-17/0 (M)	1.01/1	2.13/1	3.11/2	4.21/1	
Human (s/c)	-10/0	1.01/1	2.13/1	3.11/2	4.21/1	
Amphioxus	-17/0 (M)	1.01/1	2.13/1	3.24/0	4.21/1	
Ascidian	-10/0 (M)	1.01/1	2.13/1	3.24/0	4.21/1	
CaM						
Human (III)	-10/0 (M)	1.01/1	2.13/1	3.12/0	4.21/1	
Rat (I,III)	-10/0 (M)	1.01/1	2.13/1	3.12/0	4.21/1	
Rat (II)	-10/0 (M)	1.01/1	-	3.12/0	4.21/1	
Chicken	-10/0 (M)	1.01/1	2.13/1	3.12/0	4.21/1	
<i>Aplysia</i>	-10/0 (M)	1.01/1	2.13/1	-	4.21/1	
<i>Drosophila</i>	-10/0 (M)	-	2.13/1	-	4.21/1	
Ascidian	-10/0 (M)	1.01/1	2.04/0	3.12/0	4.21/1	
Spec						
Sea urchin	-12/0 (M)	1.01/1	2.13/1	3.18/2	4.21/1	
ELC						
Human	-09/0 (M)	1.01/1	2.12/1	3+01/0	4.21/1	
Rat (L1)	-09/0 (M)	1.01/1	2.12/1	3+01/0	4.21/1	
Rat (L4)	-08/0 (M)	1.01/1	2.12/1	3+01/0	4.21/1	
Chicken (L1)	-09/0 (M)	1.01/1	2.12/1	3+01/0	4.21/1	
Chicken (L3)	-08/0 (M)	1.01/1	2.12/1	3+01/0	4.21/1	
Parvalbumin						
Human	-	-	2.11/1	3.23/2	4.21/1	
Rat	-	-	2.11/1	3.23/2	4.21/1	
RLC						
Rat	-27/0 (M)	1.05/0	1+01/1	2+03/1	3.21/2	4.02/0

(M) shows the intron is inserted just after the initiation codon, ATG. (-) shows the absence of intron. *The last intron of *CaVP* is inserted within the 3'-noncoding region.

introns (1.01/1, 2.13/1, and 4.21/1, respectively) of *CaVP* are identical with those of TnCs, CaMs, and Spec. Thus, the gene structure of amphioxus *CaVP* is typical of the TnC superfamily and *CaVP* is clearly a member of the TnC superfamily. Within the family CaVP seems more closely related to the following three members: TnCs, CaMs, and Spec (Table II). The sixth intron, which is observed in the 3'-noncoding region, is unique to *CaVP*, and may be a characteristic feature, *i.e.* a landmark of the *CaVP* lineage.

Except for in RLC,⁴ the exon/intron organizations are well conserved among all members of the TnC superfamily (including *CaVP*). Basically they all possess five introns with the following characteristics: intron 1 is inserted just after the initiation codon (phase 0), and in a few exceptions, it is also inserted before domain I and the phase 0 mode is conserved; intron 2 is inserted at position 1.01/1 in all members without any exception; intron 3 is inserted at position 2.13/1 in *CaM*, *TnC*, *Spec*, and *CaVP*, whereas in the other members the insertion point is slightly different, but phase 1 is conserved (with the notable exception of intron 3 in the *CaMs* of the ascidian, *Halocynthia roretzi* with 2.04/0, Ref. 19); finally intron 5 is inserted at position 4.21/1 in all members without any exception.

Only the positions of intron 4 is not conserved at all in the TnC superfamily. Two explanations can account for this diversity: either intron 4 acquired different positions in the TnC superfamily genes through intron sliding, or the members of the TnC superfamily may have gained a fourth

intron independently during their evolution. According to the first hypothesis the fourth introns of the TnC superfamily are derived from a common ancestral intron, but its sequence changed so rapidly that the homology quickly disappeared. Indeed these introns are different in length and sequence; there is no positive proof that they are homologous. In general the evidence of intron sliding itself is weak (20), and besides, except in one instance, between protochordate/vertebrate *TnCs* (21), no sliding has occurred within each lineage of subfamilies. The second hypothesis implies that the ancestor gene possessed only four introns and the present-day members gained a fourth intron independently during their evolution. Surprisingly, this intron 4 was gained within or near domain III for all genes. Among the members of the TnC superfamily, the most conservative molecule is CaM, thus the common ancestor of the family is supposed to have been a CaM-like protein or, perhaps, CaM itself. Interestingly, in invertebrate (*Drosophila* and *Aplysia*) *CaM* genes (19, 22, 23) the fourth intron is absent, and this may reflect the absence of the fourth intron in the ancestor gene. The discovery of new members of the TnC superfamily may explain the evolutionary rules that govern the position of intron 4.

In this study, we elucidated the gene structure of a new Ca²⁺-binding protein and show that, although structurally and functionally it is a unique protein, it belongs to the TnC superfamily. The gene structure may be used as a tool to assign a protein to a given superfamily. This gene structure sheds more light on the evolution of constant and variable elements in the genes of the TnC superfamily.

⁴The positions of introns in the RLC are different from those of the other members of the TnC superfamily, but for conciseness this particular diversity is not analyzed in this study.

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