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Conserved sequences of sperm-activating peptide and its receptor throughout evolution, despite speciation in the sea star *Asterias amurensis* and closely related species

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Summary

The asteroidal sperm-activating peptides (asterosaps) from the egg jelly bind to their sperm receptor, a membrane-bound guanylate cyclase, on the tail to activate sperm in sea stars. Asterosaps are produced as single peptides and then cleaved into shorter peptides. Sperm activation is followed by the acrosome reaction, which is subfamily specific. In order to investigate the molecular details of the asterosap–receptor interaction, corresponding cDNAs have been cloned, sequenced and analysed from the Asteriinae subfamily including *Asterias amurensis*, *A. rubens*, *A. forbesi* and *Aphelasterias japonica*, as well as *Distolasterias nipon* from the Coscinasteriinae subfamily. Averages of 29% and 86% identity were found from the deduced amino acid sequences in asterosap and its receptor extracellular domains, respectively, across all species examined. The phylogenetic tree topology for asterosap and its receptor was similar to that of the mitochondrial cytochrome *c* oxidase subunit I. In spite of a certain homology, the amino acid sequences exhibited speciation. Conservation was found in the asterosap residues involved in disulphide bonding and proteinase-cleaving sites. Conversely, similarities were detected between potential asterosap-binding sites and the structure of the atrial natriuretic peptide receptor. Although the sperm-activating peptide and its receptor share certain common sequences, they may serve as barriers that ensure speciation in the sea star *A. amurensis* and closely related species.

Keywords: Acrosome reaction, Guanylate cyclase, Sea star, Sperm-activating peptide

Introduction

Speciation in fertilization is critical in the beginning of development. While mechanisms of fertilization have diverged from invertebrates to humans, several strictly defined steps have been governed by specific molecules from eggs and sperm for the recognition of one another (Vacquier, 1998). These steps include sperm activation and/or attraction to the egg, as well as sperm

acrosome reaction upon egg contact. Understanding the mechanisms of these molecular interactions is important, as any disruptions of these processes could cause infertility and destroy species barriers at the molecular level.

In the sea star, *Asterias amurensis*, the asteroidal sperm-activating peptide (asterosap) has been purified and identified from the extracellular jelly coat of the egg. Asterosap is composed of 34 amino acids and has an intramolecular disulphide bond that is essential for its physiological activity (Nishigaki *et al.*, 1996). The asterosap precursor mRNA encodes 10 asterosap molecules that are tandemly connected by spacer sequences (Matsumoto *et al.*, 1999). Asterosap binds to the extracellular domain of its receptor, a membrane-bound guanylate cyclase, in the sperm flagellum to activate its intracellular catalytic domain for the production of cGMP (Nishigaki *et al.*, 2000; Matsumoto *et al.*, 2003). The cooperation of three

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egg jelly coat components, ARIS (acrosome reaction-inducing substance), Co-ARIS and asterosap, induces the acrosome reaction (Hoshi *et al.*, 1994). Astrosap is essential for the induction of the acrosome reaction by elevating the internal pH of sperm (Kawase *et al.*, 2005). In addition, asterosap induces hyperactivation-like flagellar movement in sperm that are firmly attached to a solid surface (Shiba *et al.*, 2006), potentially allowing an effective acrosome reaction on the surface of the jelly coat. Furthermore, asterosap also has sperm-attracting activity (Bohmer *et al.*, 2005). The specificity of asterosap activity is conserved within the superorder level (Nakachi *et al.*, 2006).

In sea urchin, resact (SAP-IIA) from *Arbacia punctulata* is the only peptide known to have sperm-attracting activity among more than 70 sperm-activating peptides identified from the egg jelly coat (Ward *et al.*, 1985). Resact binds directly to a membrane-bound guanylate cyclase receptor in sperm and activates it (Singh *et al.*, 1988; Kaupp *et al.*, 2003).

In order to investigate the molecular details of the interaction between asterosap and its receptor, we cloned the cDNA encoding these proteins from five closely related sea star species. We report that although the amino acid sequences are highly conserved over these species, small differences in the sequences can explain their speciation. The functional mechanism of the asterosap receptor is also discussed by analogy to another membrane-bound guanylate cyclase, the atrial natriuretic peptide receptor.

Materials and methods

Animals and gametes

Sea stars were collected locally during their annual breeding seasons as described previously (Nakachi *et al.*, 2006). The sea stars used in this study were *Asterias amurensis* (Aa), *A. rubens* (Ar), *A. forbesi* (Af), *Aphelasterias japonica* (Apj) and *Distolasterias nipon* (Dn).

Cloning of genes

Total RNA was prepared from fresh testes or ovaries by the acid guanidium-phenol-chloroform (AGPC) method (Sambrook *et al.*, 1989). First-strand cDNA was synthesized using reverse transcriptase with either a random 9-mer using Rever Tra Ace (TOYOBO) or the ImProm-2 reverse transcription system (Promega) and was subsequently used as a template for PCR. DNA fragments of asterosap and spacer sequences were amplified by PCR from ovary cDNA using primers TN1 5'-ATCGCTCGAGTAAGACAA-3' and TN2 5'-TCGTTTGCTCTGACATTG-3'. Extracellular

domain sequences of the asterosap receptors were amplified by PCR from testes cDNA using primers hani1 5'-AGGTGTTTGATGCTGTCTGTCG-3', hani2 5'-GCGTAGTCTCCGTTATTGAGACC-3' MN1' 5'-AAGGAGACAACAAGGATTTAACGTG-3' and MN2 5'-GACCGGCATATCTAGAGGTGG-3'. Guanylate cyclase domain sequences of the asterosap receptors were also amplified from testes cDNA using primers P2 5'-GACATCGTCGGGTTTACC-3' and P6 5'-AGTGTTGACGGTGTCACC-3'. PCR products were separated by size via agarose gel electrophoresis, extracted and cloned into the pGEM-T vector (Promega) and sequenced with the M13 forward and reverse primers using BigDye chemistry on an ABI PRISM 3100 DNA sequencer (Applied Biosystems). We deposited the sequences into GenBank and the accession numbers are listed in the Supplemental materials section.

Sequence analysis

Multiple alignments of nucleotide and amino acid sequences were done using Clustal X (Thompson *et al.*, 1997). Phylogenetic reconstruction of nucleotide sequences was performed using PHYLIP version 3.66 (Felsenstein, 2005) and the phylogenetic trees were viewed using njplot (Perriere & Gouy, 1996). Secondary structures and β -turn structures of the amino acid sequences were predicted using PORTER (Pollastri & McLysaght, 2005) and BetaTPred2 (Kaur & Raghava, 2003), respectively. Hydropathy index and pI of amino acid sequences were calculated by sequence analysis (Gilbert, 2006) using Kyte & Doolittle parameters (Kyte & Doolittle, 1982) in a running average of five amino acids. The coiled-coil structures in the amino acid sequences were predicted by COILS (Lupas *et al.*, 1991). Accession numbers for the guanylate cyclase genes were X12874 from sea urchin (*Arbacia punctulata*; Ap) and AB070354 from Aa. Accession numbers for the mitochondrial cytochrome c oxidase 1 sequences other than ours were AF498647 (Af), AF031028 (Ap) and AF525453 (*Hemicentrotus pulcherrimus*, Hp, sea urchin). Sequence alignments referring to their 3D structures were performed by the 3D-PSSM (Kelley *et al.*, 2000) and Swiss model (Schwede *et al.*, 2003) servers. The hormone-bound, atrial natriuretic peptide receptor extracellular domain (Ogawa *et al.*, 2004) was accessed from the Protein Data Bank (ID code 1t34).

Results

In order to investigate the molecular details of the interaction between asterosap and its receptor, we cloned the cDNAs encoding these proteins from five closely related sea star species.

Sequence variation among asterosap

Sea star asterosap sequences exist as 10 consecutive repeats interspersed with short spacers (Matsumoto *et al.*, 1999). These 10 asterosap sequences from the same genome are not identical but vary among them. The cDNAs of randomly chosen seven asterosap isoforms were each cloned including their spacer regions from five sea star species: *Asterias amurensis* (Aa), *A. rubens* (Ar), *A. forbesi* (Af) and *Aphelasterias japonica* (Apj), from the Asteriinae subfamily and *Distolasterias nipon* (Dn) from the Coscinasteriinae subfamily. All these five species belong to the family Asteriidae.

The deduced amino acid sequences of the asterosaps were highly conserved among the species with more variation found in the N-terminal region (Fig. 1). Of the 35 cloned sequences, the asterosaps had 10 completely conserved amino acid sites out of 34, covering 29% of the peptide length. While all asterosaps were of the same length, the spacer regions varied ranging from 12 to 22 amino acids. The spacer regions had four completely conserved amino acids sites out of 12 (excluding gap sites), corresponding to 33% identity.

In *A. amurensis* (Aa), asterosap has an intramolecular disulphide bond between Cys8 and Cys32, which is known to be essential for its activity (Nishigaki *et al.*, 1996). These Cys residues were among the completely conserved amino acids from all asterosap sequences examined. As shown in Fig. 1, some amino acid substitutions were unique in *Asterias*, in Apj or in Dn. All the spacer regions had paired basic residues, Lys–Arg or Lys–Lys, at both ends, which are commonly found at the cleavage sites of pro-hormone precursors (Hook *et al.*, 2004). The results suggest that the asterosap precursor undergoes endoproteolytic processing at either the C- or N-termini of paired basic residues followed by the removal of the remaining basic residues as shown in other pro-hormone processing pathways (Fricker, 1988; Steiner, 1998; Hook *et al.*, 2004). The pI represents the favourable environmental pH range of biologically active molecules such as asterosap. While the distribution of pIs for asterosap was between 4.5 and 10.0 in Aa, Af, Ar and Apj, pIs for asterosaps from Dn range from 3.8 to 6.3.

Every asterosap was predicted to have two amphipathic helices between two conserved Cys residues (Fig. 1). Positive charges (Arg11 and 13) in the first helix and negative charges (Asp20 and Glu21) in the second helix were highly conserved. Then, we assumed that these helical structures are important for asterosap binding to its receptor. Hydropathy index of the asterosaps indicated the overall hydrophilicity, explaining its diffusive characteristic in seawater. Conserved hydrophobic residues in each helix may also

be involved in either binding to its receptor or peptide folding. Spacer regions had turn structures and were entirely hydrophilic.

Asterosap receptor

A. amurensis asterosap receptor has four domains: extracellular, transmembrane, kinase-like and guanylate cyclase (Matsumoto *et al.*, 2003). We cloned the cDNAs of the extracellular and guanylate cyclase domains of the asterosap receptor from four additional sea star species and compared these sequences. The deduced amino acid sequences (Fig. 2) shared high similarities in these domains: 86% in the extracellular domain and 94% in the guanylate cyclase domain. Most amino acid differences were found between *Asterias* (Aa, Ar and Af) compared with Apj and Dn: more than 10% of the extracellular domain sequences differed. In the extracellular domain, all four Cys residues and three potential glycosylation sites were conserved among the five species. In the guanylate cyclase domain, all putative catalytic sites (Liu *et al.*, 1997) were conserved (Fig. 2).

Molecular evolution of asterosap and its receptor

In order to compare the evolutionary patterns between asterosap and its receptor sequences, we performed phylogenetic reconstruction at the nucleotide level (Fig. 3). The phylogenetic tree reconstructed using the 35 asterosap sequences showed three major groups, each corresponding to the three subfamilies: Dn, Apj and *Asterias*, although the bootstrap analysis did not support these three clusters (Fig. 3a). Interestingly, clustering among *Asterias* asterosap sequences was not consistent with species clustering. For example, the clusters from a to d in Fig. 3a included more than two *Asterias* species. Moreover, identical asterosap sequences were found between Af02 and Aa13 and between Apj07 and Af05. More phylogenetics analysis may be required to establish taxonomical relationships among these sea star species, especially the phylogenetic location of Apj. The occurrence of close relationships among *Asterias* species found in phylogenies agrees with our previous results on the gamete-compatibility grouping in sea stars (Nakachi *et al.*, 2006).

Discussion

Interaction between asterosap and its receptor

The structure of the asterosap receptor and the binding of the asterosap to its receptor were modelled based on the structure of the rat atrial natriuretic peptide and its receptor (ANP and NPR-A; PDB entry 1t34).

Figure 1 Comparison of asterosap and spacer amino acid sequences from five sea star species. Plus (+) and minus (–) position numbers at the top indicate positions within the spacers from the N- and C-termini, respectively. A dot (·) shows the residue identical to Aa01; a minus sign (–) indicates a gap. The slash (/) in the ruler shows the potential cleavage site. Sequences used for PCR primers are underlined by dotted lines in Aa01 and primer names are indicated in square brackets ([]) above with arrows indicating the amplification directions. Two conserved Cys residues are marked by number signs (#) above Aa01. Positively and negatively charged conserved residues in asterosaps are shaded with light and dark grey, respectively. The ‘Common’ row shows the symbols based on the sequence similarity for each position after CLUSTAL X: an asterisk (*) for a completely conserved position; a colon (:) for the position conserved among the ‘strong’ amino acid groups; a period (.) for the position conserved among the ‘weaker’ amino acid groups. The ‘Motif’ row shows the predicted secondary structures for Aa04: a tilde (~) for the coil structure; and an equal (=) for the helical structure. The ‘Phobic’ row shows Kyte & Doolittle hydrophathy index from a running average of five amino acids: a plus (+) for hydrophobicity; a vertical bar (|) for hydrophilicity.

receptor in sperm appears to be present as a dimer and oligomer (data not shown). In addition, the Aa asterosap receptor has a potential coiled-coil region in the area similar to that of NPR-A, which is important for dimerization (Wilson & Chinkers, 1995). A sequence

		{hani1⇒}	
Aa	-20	-----MRCIMLSVV-----	-12
		+1	
		Signal ← →Extracellular (+++)	
Aa	-11	LVAGYVWVALGTNFKIGLLVPLTDPQTGNASGFGDPVAGAFPVAVDDINLNPAILPGHTV	49
Ar	-11		49
Af	-11	F.....D.....	49
Apj	-11	EY.....N.....I.....M.....I	49
Dn	-11	EY.....N.....I.....L	49
		:*****:*****:*****:*****:*****:*****:*****:	
		# # #	
Aa	50	SWEWVDTKCDINTGLTAVSDWWKRGFVGVI GPGCSCDYEARLAGSINFPMDYGCDEGAV	109
Ar	50		109
Af	50	G	109
Apj	50	S.....	109
Dn	50	S.....	109
		*****:*****:*****:*****:*****:*****:*****:	
		(+++)	
Aa	110	SNKLLYPTYIRTLPSPSTRIVDALIVTLQKFDWDQVTVVYRNHSIWNTNINAMKEEFVH	169
Ar	110	F.....S.....	169
Af	110		169
Apj	110	V.....SV·I··M.....V·N··YY··R··NK	169
Dn	110	V.....SV·I··M.....V··YY··R··NK	169
		*****:*****:*****:*****:*****:*****:*****:	
		(+++)	
		{MN1'⇒}	
		{hani2⇐}	
Aa	170	ITVOHOFVFPOTGFPNNDISIINPFPEIFTRTKETTRIVVFLGEMIELRSFAMAALDEGLN	229
Ar	170		229
Af	170	E.....	229
Apj	170	E·A.....A.....T.....I.....L.....V.....Q.....	229
Dn	170	E.....A.....V.....I.....L.....V.....V.....Q.....	229
		*****:*****:*****:*****:*****:*****:*****:	
Aa	230	NGDYAILGMAIDHKIRRSQNWHS�FLHMGTYLDEKAAKAMESVLI IAPKAPKFTFVYKS	289
Ar	230	L.....T.....	289
Af	230		289
Apj	230	IA.....NT.....Y.....L.....	289
Dn	230	IA.....S.....Y.....V.....L.....	289
		*****:*****:*****:*****:*****:*****:*****:	
Aa	290	WNVKVRDSVQGAPFFQTGREFHTFSAFLYDATILFAKALEETLAAGEDPFDGEAIVSHAM	349
Ar	290		349
Af	290		349
Apj	290	N·KA.....A.....AA.....S.....	349
Dn	290	·I··NR·EE.....A.....AD.....S.....	349
		;*: *: *****:*****:*****:*****:*****:*****:	
Aa	350	GVQYQSISMLQNGIDESGDGISRYMLMDMNELQEADSWLTAGYPGVIGVGEFIRNSNGRW	409
Ar	350	L.....	409
Af	350		409
Apj	350	·T.....E·D.....I.....S·QG.....Q.....	409
Dn	350	·T.....E·VY.....P·KG.....H.....	409
		*.****** ** *****:*****:*****:*****:*****:*****:	

Figure 2 Comparison of asterosap receptor amino acid sequences from five sea star species. A dot (·) indicates the amino acid identical to Aa at the corresponding position. Mature proteins start from the residue +1 after the N-terminal signal peptides. Each domain starts at |→ and ends at ←|. Sequences used for PCR primers are underlined with dotted lines in Aa and primer names are given in square brackets ([]) with arrows indicating the amplification directions. An asterisk (*), a colon (:), and a period (.) at the bottom show the levels of sequence similarities for each position (see the Fig. 1 legend). Extracellular Cys residues are marked by a number signs (#). Potential N-glycosylation sites (Asn-X-Ser/Thr) are marked by a plus (+) in parentheses. Parts of the Aa sequence subjected to the homology modelling are underlined. A potential ligand-binding region is double underlined with positively and negatively charged residues shaded with light and dark grey, respectively. Potential phosphorylation sites in the region equivalent to the rat atrial natriuretic peptide receptor are shaded with light grey in Aa (from positions 500 to 517). A potential coiled-coil region is underlined by short- and long-dotted lines. Potential GTP binding sites in the guanylate cyclase domain are marked by 'g'. Except for Aa, only extracellular and guanylate cyclase domains were cloned and sequenced.

Figure 2 (Cont.)

Resact is a sperm-activating peptide from *Arbacia punctulata* (Ap) and is composed of 14 residues with

an intramolecular disulphide bond. The primary structure of the resact receptor was also compared with the astrosap receptor to predict their functional similarity. Their extracellular domains have high sequence similarity. While the resact receptor lacks the large C-terminal portion of the guanylate cyclase domain including almost all putative key catalytic residues, the astrosap receptor and resact receptor may have similar modes of ligand binding but different catalytic regulatory mechanisms.

Figure 3 Phylogenetic relationships reconstructed based on the nucleotide sequences of asterosap (a) and its receptor (b and c). The receptor phylogenies were reconstructed using the extracellular domain (B, 1299 nucleotides) and the guanylate cyclase domain (C, 326 nucleotides) separately. Neighbour-joining phylogenetic method (Saitou and Nei, 1987) was used on nucleotide distances estimated by F84 model (Kishino and Hasegawa, 1989). Bootstrap analysis was done with 1000 pseudoreplicates and supporting values greater than 80% are shown on each consensus phylogeny. For asterosap receptor analyses, resact receptor sequences from sea urchin were added as outgroups: Ap, *Arbacia punctulata*; Hp, *Hemicentrotus pulcherrimus*.

Survival strategies of sea stars

The acrosome reaction is an important fertilization process in sea stars, which takes place upon the arrival of sperm onto the egg jelly surface. ARIS, a huge proteoglycan, is the major component of egg jelly in sea stars. A sulphate moiety in ARIS plays an important role in the acrosome reaction. The pH of the egg jelly and the neighbouring seawater can be affected by the content of the sulphate moiety caused by its acidic properties. Each species has several varieties of asterosaps at different pIs. This variation is advantageous for sea stars because a wide distribution of pIs allows with the organism to adjust to any environmental changes. Alternatively, differences in pI can reflect the different sugar components of egg jelly in

sea stars. For example, Dn has fewer sulphate moieties in its egg jelly, which results in a different pH in the environment.

Aa, Apj and Dn live in close proximity off the Japanese shore in the North Pacific Ocean. There is, therefore, a certain risk of cross-fertilization between them. The fertilization season of Apj (fall) is different from that of Aa and Dn (early spring). While Aa and Apj are able to induce cross-acrosome reactions *in vitro* using each other's egg jelly (Nakachi *et al.*, 2006), natural cross-fertilization is prohibited by the season. The cross-acrosome reaction (i.e. fertilization) is, therefore, unlikely to occur between Aa, Apj and Dn. These facts are evident in the phylogenetic relationship presented in Fig. 3.

Figure 4 A schematic model of hormone and receptor binding. The rat atrial natriuretic peptide (ANP) and its receptor (NPR-A) are shown on the left. Asterosap and its receptor are shown on the right. Two identical receptor subunits are connected by ionic interactions at the top. A large hydrophobic tryptophan (W) residue from each subunit acts as a spring to secure the hinge. Interactions between the negatively charged aspartic acid (D) and the positively charged arginine (R) play important roles in the binding between ligands and the receptors. The molecular size of asterosap peptide (3700 Da, 34 amino acids) is larger than that of ANP (3100 Da, 28 amino acids).

We have identified homologous guanylate cyclase signalling systems from five different sea star species, presumably the first attempt in marine invertebrates from the North and South Pacific Ocean as well as the Atlantic Ocean. The cloned asterosap and guanylate cyclase sequences both exhibited speciation. Although the number of clones was limited caused by specimen availability, the analysed codon utilization did not have a strong bias. As we accumulate more clones, we will have the capacity to test evolutionary hypotheses, such as positive selection and co-evolution of asterosap and its receptor.

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