RESEARCH COMMUNICATION

The species-specific egg receptor for sea urchin sperm adhesion is EBR1, a novel ADAMTS protein

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Species-specific adhesion of sperm to the egg during sea urchin fertilization involves the interaction of the sperm adhesive protein, bindin, and a complementary receptor on the egg surface, and serves to restrict the gene pool to individuals of the same species. We used PCR representation difference analysis to clone the species-specific egg receptor for bindin, EBR1, from Strongylocentrotus franciscanus (Sf) and S. purpuratus (Sp). Sf-EBR1 contains a novel ADAMTS-like N-terminal domain followed by ~19 tandem EBR repeats consisting of alternating CUB and thrombospondin type 1 (TSP-1) domains where the last 10 EBR repeats are species-specific and highly conserved. Recombinant protein corresponding to the species-specific EBR repeat displays species-specific sperm adhesion and bindin-binding activity. The Sp-EBR1 ortholog has the same ADAMTS (a disintegrin and metalloprotease with thrombospondin type-1 modules) core region followed by eight and one-half tandem egg bindin receptor (EBR) repeats that share 88% identity with the Sf-EBR1 repeats, but has an entirely different species-specific domain consisting of hyalin-like (HYR) repeats. Thus, the species-specific domains of egg bindin receptor 1 (EBR1) from both species function as the egg surface receptor to mediate species-specific sperm adhesion.

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Sperm–egg interaction typically exhibits species specificity in most animals. Several molecules from both sperm and eggs that are thought to mediate gamete interaction have been identified in many animals. In abalone, the acrosomal protein of sperm, lysin, interacts with the egg vitelline envelope receptor for lysin (VERL) and creates a hole on the extracellular matrix (ECM) of the egg with species selectivity (Swanson and Vacquier 1997). In mammals, numerous molecules, such as galactosyltransferase, ADAM (a disintegrin and metalloprotease), and Sp56, have been identified from sperm and are proposed to interact with such egg zona pellucida proteins Zp2, Zp3, or Integrins on the egg. However, a clear understanding of the specificity of these interactions and the regulation of species specificity is not yet known (Primakoff and Myles 2002; He et al. 2003).

Acrosome-reacted sea urchin sperm adhere to glycoprotein receptors that are components of the vitelline envelope (VE) known as the ECM of the egg. It has been known for several decades that the protein bindin on the acrosome-reacted sea urchin sperm head mediates the species-specific adhesion of sperm to the VE during fertilization (Summers and Hylander 1975; Glabe and Vacquier 1977b; Vacquier and Moy 1977; Glabe et al. 1982). The egg ECM also serves as a major barrier for interspecific fertilization in mammalian systems (Yanagimachi 1981, 1994). In sea urchins, the complementary receptor for bindin was shown to be an ~350-kD protein on the VE of the egg (Glabe and Vacquier 1978; Carroll et al. 1986; Ruiz-Bravo and Lennarz 1986; Ohlendieck et al. 1993). However, the identity of this receptor has remained elusive, and attempts to solubilize and purify the active receptor from the VE of the egg have not been successful (Glabe and Lennarz 1981, Ruiz-Bravo et al. 1986). Therefore, we used a novel molecular approach, representation difference analysis (RDA), to amplify species-specific cDNA fragments from ovaries of the sea urchin Strongylocentrotus franciscanus (Sf) that are distinct from cDNAs fragments in another species of the same genus, S. purpuratus (Sp), to identify genes involved in determining species specificity (Hubank and Schatz 1994, Kamei et al. 2000).

Here we report the identification and characterization of the species-specific egg bindin receptor (EBR1) from two sea urchin species, S. franciscanus and S. purpuratus, along with their unique domain structures. The biological activity of recombinant species-specific egg bindin receptor [EBR] and hyalin-like [HYR] repeats indicate that they serve as species-specific ligands for the sperm adhesive protein bindin.

Results and Discussion

Characterization of the species-specific cDNA clone, Sf-H2

We isolated four species-specific cDNA clones from S. franciscanus ovary mRNA, as previously described (Kamei et al. 2000). One of the clones (Sf-H2) was a candidate for the egg-bindin receptor because it hybridizes species-specifically to an ~14–15-kb mRNA (see below) and therefore is sufficiently large to encode the 350-kD egg receptor protein. We raised polyclonal antibodies against recombinant Sf-H2 protein, and the antibodies detect a protein of ~350 kD on purified VE s from S. franciscanus, but not S. purpuratus eggs (Fig. 1A,B). Thus, the Sf-H2 cDNA probe and antibody detect molecules of the size predicted for the egg receptor.

Because the Sf-H2 transcript encodes a species-specific VE protein of the size and location expected for the receptor, we determined whether Sf-H2 displayed species-specific adhesive activity by using several different assays. We first examined the effect of anti-Sf-H2 antibody or recombinant Sf-H2 protein on fertilization. Anti-Sf-H2 polyclonal antibody inhibited fertilization of S. franciscanus eggs in a concentration-dependent manner, but
had no effect on the fertilization of *S. purpuratus* eggs [Fig. 2A]. Similarly, recombinant *Sf*-H2 protein only inhibited *S. franciscanus* fertilization, but not *S. purpuratus* fertilization [Fig. 2B]. These results suggest that anti-*Sf*-H2 antibodies block the receptor on the egg, and the recombinant *Sf*-H2 protein competitively inhibits the binding of sperm to the VE receptor. We also directly examined sperm adhesive activity of *Sf*-H2 by using a recombinant glutathione S transferase (GST) fusion protein immobilized on glutathione agarose beads to assess sperm adhesion to the beads. We found that acrosome-reacted *S. franciscanus* sperm specifically bound to beads containing immobilized *Sf*-H2 GST fusion protein, but not to control beads containing GST alone [Fig. 2C]. These data are consistent with the interpretation that *Sf*-H2 encodes a species-specific sperm receptor.

The sperm adhesive protein bindin is commonly purified as insoluble acrosomal granules, and these bindin granules were shown to species-specifically agglutinate eggs of the same species (Glabe and Lennarz 1979). This agglutination activity of bindin granules can be used to measure the adhesive interaction between bindin and the egg surface (Glabe and Lennarz 1979). Recombinant *Sf*-H2 protein specifically inhibits agglutination of *S. franciscanus* eggs by homologous bindin, but does not inhibit the agglutination of *S. purpuratus* eggs by *S. purpuratus* bindin [Fig. 2D]. We also directly measured the interaction of iodinated recombinant *Sf*-H2 protein to purified bindin granules using a sedimentation assay. Recombinant *Sf*-H2 protein bound only to *S. franciscanus* bindin granules in a concentration-dependent and saturable manner [Fig. 2E]. Therefore, *Sf*-H2 satisfies a number of different functional criteria for the egg surface adhesive receptor for bindin.

The egg surface receptor for sperm is either released from the VE or destroyed due to the action of proteases that are released from the cortical granules upon fertilization (Vacquier et al. 1973; Glabe and Vacquier 1977a). We examined the localization of the *Sf*-H2 on whole mounts of fertilized and unfertilized eggs by immunofluorescence staining with anti-*Sf*-H2 antibodies. Bright staining was detected on the unfertilized egg surface, but

![Figure 1](image1.png)  
**Figure 1.** Species specificity of *Sf*-H2. (A) Coomassie blue staining of purified VEg from *S. franciscanus* and *S. purpuratus* separated on a 4%–20% SDS-PAGE gel. [Lanes 1–3] *S. franciscanus* VE. [Lanes 4–6] *S. purpuratus* VE. (B) Western blot analysis of VE from *S. franciscanus* and *S. purpuratus* separated on a 4%–20% SDS-PAGE gel (shown in A) using anti-*Sf*-H2 antibody. The arrow highlights the ~350-kD protein detected by the anti-*Sf*-H2 antibody. [Lanes 1–3] *S. franciscanus* VE. [Lanes 4–6] *S. purpuratus* VE. Lanes 1 and 4 are 2 μg of VE protein, lanes 2 and 5 are 5 μg of VE protein, and lanes 3 and 6 are 10 μg of VE protein.

![Figure 2](image2.png)  
**Figure 2.** Inhibition of fertilization by anti-*Sf*-H2 antibody and recombinant *Sf*-H2 protein. (A) Species-specific inhibition of *S. franciscanus* fertilization by anti-*Sf*-H2 antibody. Increasing concentrations of the antibody only inhibit the fertilization of homologous eggs. Red line indicates *S. franciscanus* gametes, black line indicates *S. purpuratus* gametes. (B) Inhibition of fertilization by recombinant *Sf*-H2. Increasing concentrations of *Sf*-H2 inhibits only the fertilization of homologous eggs. Both red and blue lines show the percentage of inhibition of fertilization by using recombinant *Sf*-H2–GST. Both broken lines indicate control samples GST only. (C) Species-specific binding of acrosome-reacted sperm to immobilized recombinant *Sf*-H2 protein. The graph shows the number of sperm binding to the beads. Red line indicates *S. franciscanus* sperm; blue line, *S. purpuratus* sperm. Both broken lines show the binding to GST controls. (D) Species-specific inhibition of bindin-mediated egg agglutination by recombinant *Sf*-H2 protein. The percentage of agglutinated eggs is shown as a function of increasing recombinant *Sf*-H2 protein. Red line indicates *S. franciscanus* eggs and *S. franciscanus* bindin; black line indicates *S. purpuratus* eggs and *S. purpuratus* bindin. (E) Species-specific binding of recombinant iodinated *Sf*-H2 to purified bindin particles. Red line indicates *S. franciscanus* bindin; black line, *S. purpuratus* bindin. (F) Immunostaining of unfertilized and fertilized *S. franciscanus* eggs with anti-*Sf*-H2 antibody. Panel 1 shows the phase contrast image; panel 2, the bright-field immunofluorescence staining with anti-*Sf*-H2 antibody. The egg on the left is unfertilized, and the one on the right is fertilized.
this immunoreactivity was greatly diminished on the fertilization envelope and egg surface after fertilization (Fig. 2F). The absence of anti-Sf-H2 staining in the fertilization envelope of fertilized eggs is consistent with the known release or destruction of the receptor from the elevating fertilization envelope after fertilization.

The Sf-H2 gene product meets all of the known structural and functional criteria for the egg receptor, including size, VE localization and modification after fertilization, species-specific sperm, and bindin-binding activity. Thus, from these data, we conclude that Sf-H2 is the egg receptor for sperm, and we have named this gene product egg bindin receptor (EBR1).

**Comparison of the sequences of EBR1 from both S. franciscanus and S. purpuratus**

The deduced amino acid sequence of Sf-EBR1 consists of 4595 amino acids (GenBank accession no. AY138966). Starting at the N terminus, the protein contains a 30-residue signal peptide followed by a putative propeptide, metalloprotease domain, cis-rich domain, and eight thrombospondin type 1 (TSP-1) domains, indicating that EBR1 shares several significant structural features with members of the ADAMTS family of proteins (a disintegrin and metalloprotease with thrombospondin type-1 modules; Tang and Hong 1999; Tang 2001). However, Sf-EBR1 lacks a disintegrin-like domain that typically follows the catalytic domain in ADAMTS family members. Sf-EBR1 also contains an additional C-terminal region containing ~19 repeats of a novel motif (EBR repeats) located immediately after the ADAMTS-like core sequence (Fig. 3A). These novel motifs contain a TSP-1 and a CUB (C1s/C1r, uEGF and bone morphogenic protein) subdomain (Bork and Beckmann 1993; Adams 1997). Each TSP-1 domain is from 53 to 59 amino acids, and each CUB domain is ~104–123 amino acid residues.

Thus, each EBR repeat is ~513 bp or 171 amino acids. The nine EBR repeats that follow immediately after the ADAMTS-like core region of Sf-EBR1 are significantly more variable than the last 10 EBR repeats and are defined as core EBR repeats because they also occur in Sp-EBR1 (TSP, thin light blue ovals; CUB, yellow ovals; Fig. 3A). The last 10 EBR repeats of Sf-EBR1 are homogeneous (TSP, thin green ovals; CUB, dark blue ovals), and each domain shows 85%–100% identity with the other repeats in this region. The last 10 homogeneous repeats are also species-specific, because they do not hybridize to S. purpuratus mRNA and are absent in Sp-EBR1 (see below). The original cDNA fragment isolated by RDA, Sf-H2, corresponds to one of these species-specific EBR repeats. In contrast, the TSP-1 domains outside of the EBR repeats in the ADAMTS-like core region and the TSP-1 subdomains present in the first nine core EBR repeats (thin light blue ovals) contain only 12%–50% identity with the TSP-1 subdomains in the species-specific EBR repeats (green ovals). Similarly, the first nine and the last three CUB subdomains that are in the EBR repeat (yellow ovals) display only 11%–20% identity with the CUB subdomains in the species-specific EBR repeats (dark blue ovals; Fig. 3A). The second core EBR repeat for S. franciscanus (also for S. purpuratus; see Fig. 3A) has a duplicated CUB domain in contrast with all other EBR repeats, which have one CUB domain.

Although the species-specific Sf-EBR1 repeats do not hybridize with S. purpuratus DNA, S. purpuratus has a homologous EBR1 gene because the 3’ UTR of Sf-EBR1 hybridizes to the same size fragment on genomic Southern blots of S. purpuratus DNA [data not shown]. The EBR1 gene was cloned and sequenced from an S. purpuratus ovary cDNA library by using the 3’ UTR of Sf-EBR1 as an initial probe and the same cloning and sequencing strategy used for Sf-EBR1 [GenBank accession no. AY341264]. The gene encodes a protein with a strik-
ingly similar repeat structure but with some notable differences compared with Sf-EBR1. The deduced amino acid sequence of Sp-EBR1 consists of 3713 amino acid residues. Starting at the N terminus, the protein contains a 30-residue signal peptide followed by a putative propeptide, metalloprotease domain, cis-rich domain, eight TSP-1 domains, and eight and one-half core EBR repeats. Thus, Sp-EBR1 shares the same domain structure as Sf-EBR1 for the first ~8 kb of sequence and shares 88% nucleotide identity with Sf-EBR1 in this region [Fig. 3B]. This includes the first eight and one-half core EBR repeats, which have 80%–90% identity with the corresponding EBR repeats in Sf-EBR1. The last CUB domain of both Sp-EBR1 and Sf-EBR1 also shares 86% identity [Fig. 3B]. However, the region of Sp-EBR1 corresponding to the species-specific Sf-EBR1 domains [dark blue and green ovals or white ovals] contains 11 repeats of a HYR domain [purple boxes or white boxes] and not the homogeneous EBR repeats observed in Sf-EBR1 [Fig. 3A,B]. Each HYR repeat is 243 bp or 81 amino acids, and each repeat, except the first and the last HYR repeat, shows ~95% sequence identity. The HYR repeat domain in Sp-EBR1 is distinct from the previously reported hyalin gene containing 12 HYR repeats in S. purpuratus [Wessel et al. 1998], with <40% amino acid identity when compared with HYR-like domains in Sp-EBR1. We examined the adhesive function of the Sp-EBR1 by measuring the adhesion of sperm to recombinant Sp-HYR expressed as a GST fusion protein. S. purpuratus sperm bind specifically to Sp-HYR compared with S. franciscanus sperm and GST controls [Fig. 4]. Therefore, the species-unique HYR domain of Sp-EBR1 functions as a species-specific receptor for sperm adhesion analogous to the species-specific EBR repeats of Sf-EBR1 [Fig. 2C]. The deduced EBR1 sequences from both species lack an identifiable transmembrane domain, but EBR1 would be expected to be incorporated into the VE ECM based on its similarity to other ADAMTS proteins [Kuno and Matsushima 1998].

Northern blot analyses were performed with both common and species-specific probes to examine if the probes hybridize to the same size transcript. The common probe derived from the 5′ region of Sf-EBR1 detected a single mRNA species of ~14–15 kb for S. franciscanus and 12–13 kb for S. purpuratus. Each species-specific region of EBR1 hybridized to the same size transcript identified with the 5′ common region probe, but in a species-specific manner [Fig. 5]. Reverse transcriptase PCR (RT–PCR) analysis also showed that the EBR1 gene that was originally cloned from ovary cDNA library exists in unfertilized eggs but is not detected in sperm [data not shown].

We have cloned and sequenced the gene encoding the species-specific egg bindin receptor, EBR1, from both S. franciscanus and S. purpuratus. These genes are novel ADAMTS-like proteins and share a long repetitive architecture with species-specific domains unique to S. franciscanus and S. purpuratus that are situated near the 3′ end. We have also demonstrated that the species-unique EBR1 repeats of S. franciscanus and S. purpuratus display species-specific adhesive activity by using a variety of different assays. The finding that recombinant EBR1 protein displays species-specific sperm adhesion and bindin-binding activity indicates that protein, but not carbohydrate, is important for determining the species specificity of sperm–egg adhesion. Based on the lack of an identifiable transmembrane domain and the similarity to ADAMTS, it is likely that EBR1 is an ECM protein, which is consistent with the fact that sperm bind to the VE, which represents the ECM of the egg (Glabe and Vacquier 1978; Hirohashi and Lennarz 1998). The repeated structure of the adhesive domain may be important for determining the strength of the adhesive interaction by providing multivalency. This correlates well with the structure of bindin, which has a complementary repeated structure [Minor et al. 1991].

EBR1 also has a variant ADAMTS domain at its 5′ end consisting of a signal peptide, propeptide domain, and zinc metalloprotease domain, but it lacks a disintegrin-like domain. Instead, an ADAM cysteine-rich [ACR] domain, which is typical of ADAM and snake venom proteases, as well as ADAMTS, immediately follows the zinc metalloprotease domain. The zinc metalloprotease domain contains a catalytic site consensus sequence identified in snake venom and matrix metalloproteinases [Bode et al. 1993]. This suggests that EBR1 may have protease activity, but further study is required to determine whether this is the case. Although EBR1 lacks a disintegrin-like domain, other ADAMTS family members, such as ADAMTS1 and ADAMTS5, also lack disintegrin domains [Adams and Tucker 2000]. Like other ADAMTS members, EBR1 has TSP-1 repeats after the ACR domain. The TSP-1 repeats are found in "matricellular" proteins [Bornstein 2001] and are believed to be
involved in a variety of cellular processes, including cell and matrix adhesion, cell migration, and sulfated polysaccharide binding. TSP-1 repeats fold into a novel, antiparallel, three-stranded fold capped by disulfide bonds on each end [Tan et al. 2002]. After the TSP-1 repeats of EBR1, there are a series of alternating CUB and TSP-1 repeats. Although CUB and TSP-1 domains occur together in other proteins, this is the first instance in which they have been found in a tandemly repeated fashion that we refer to as an EBR repeat. Both the CUB subdomain of the EBR repeat and the HYR domains are predicted to fold as immunoglobulin-like β-sandwich structures (Bork and Beckmann 1993; Callebaut et al. 2000). The CUB domain protein, spermadhesin, has been implicated in mammalian sperm adhesion by binding to the zona pellucida (Romero et al. 1997). The known adhesins implicated in mammalian sperm adhesion by binding to structures (Bork and Beckmann 1993; Callebaut et al. 1997) with glutathione agarose beads loaded with GSTfusionprotein, samples were inseminated with a concentration of sperm and 
S. franciscanus

Animal collection

Sea urchins, S. franciscanus and S. purpuratus, were obtained from Marinus, Inc.

Generation of Sf-H2 polyclonal antibody

Polyclonal antibodies were produced in female rabbits (New Zealand white) by subcutaneous injection of purified Sf-H2 GST fusion protein. Anti-Sf-H2 antibodies were purified with protein A Sepharose column chromatography and then applied on a GST column to remove anti-GST antibodies.

Northern blot analysis

Northern blot analysis was performed with isolated S. franciscanus and S. purpuratus ovary mRNA probed with 32P-labeled cDNA of Sf-EBR1, and Sp-HYR domain DNA (Kamei et al. 2000).

Western blot analysis

VEs were prepared from both species of sea urchin eggs (Glabe and Vacquier 1977a; Hirohashi and Lennarz 1998) and separated by electrophoresis on 4–20% SDS-PAGE gradient gels. Each sample was loaded at 2, 5, and 10 µg per lane and detected with 1:2500 dilution of anti-Sf-H2 polyclonal antiserum as the primary and 1:25,000 dilution of HRP conjugated rabbit F(ab')2 antibody as the secondary antibody.

Inhibition of fertilization

The inhibition of fertilization assay was performed with S. franciscanus and S. purpuratus eggs fertilized with sperm of the same species. After a 10-min preincubation with each concentration of antibody or recombinant protein, samples were inseminated with a concentration of sperm adjusted to give ~70–80% fertilization in the absence of inhibitor. The fertilization reaction was stopped after 5 min by the dilution of the sperm with filtered seawater. The fertilized eggs were scored by counting with a stereo microscope [Minor et al. 1993].

Sperm–beads adhesion assay

The sperm-binding assay was performed as previously described (Stears and Lennarz 1997) with glutathione agarose beads loaded with Sf-H2 GST fusion protein. GST fusion protein alone was used as a control. The numbers of attached sperm to at least 30 beads were counted by observation with a microscope.

Egg agglutination assay

Egg agglutination assays were performed as previously described (Glabe and Vacquier 1977b; Glabe and Lennarz 1979) by using recombinant Sf-H2 protein as an inhibitor. Agglutination of both species of eggs by homologous bindin was performed with increasing concentrations (0–40 µg/mL) of recombinant Sf-H2 and a constant concentration of bindin, which gave at least 75% egg agglutination.

Receptor ligand binding using iodinated EBR1

Recombinant 125I-labeled Sf-H2 protein was iodinated by using iodogen (Pierce) according to the manufacturer’s instructions and was incubated with purified bindin particles for 10 min in filtered seawater-Tris (pH 7.5) at room temperature. The sample was layered on a 20% sucrose pad in a 0.4 mL Eppendorf tube and centrifuged at 15,000 × g for 10 min to separate the iodinated protein bound to bindin particles from the unbound protein. The pellet and supernatant fractions were counted by using a γ counter. The nonspecific background was determined by adding an excess of cold recombinant Sf-H2 protein.

Localization of receptor protein on the egg

For antibody staining, 2.5% paraformaldehyde/1% glutaraldehyde-fixed unfertilized and fertilized eggs were used. Fixed eggs were washed with Tris-buffered saline (TBS; pH 7.5) several times, and then blocked with 1% BSA in TBS. The samples were incubated with anti-Sf-H2 antiserum at a 1:10 dilution for 1 h, washed three times with 0.1% Tween-TBS (pH 7.5), and then placed in blocking solution a second time. The TRITC-labeled goat F(ab')2, antirabbit F(ab')2, antibody was treated at a 1:1000 dilution for 1 h and washed with 0.1% Tween-TBS (pH 7.5) thoroughly, and samples examined under the fluorescence microscope.

cDNA library screening and isolation of the complete cDNA

S. franciscanus cDNA libraries from sea urchin ovaries were constructed by using λ ZAP (Stratagene) with oligo dT primer or random primer as recommended by the manufacturer. The S. purpuratus library was kindly provided by Dr. B. Wesel (Brown University, Providence, RI). The library was screened by using 32P-labeled Sf-H2 or unique 5’ and 3’ DNA fragments as probes. The complete gene sequence of the cDNAs was obtained by cDNA primer walking and the determination of 5’ end of cDNA by the RACE (rapid amplification cDNA ends) analysis for both S. franciscanus and S. purpuratus. The open reading frame and domain structure was assembled and analyzed by using Lasergene analysis software (DNASTAR Inc; Burland 2000) and SMART search (Letunic et al. 2002).

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