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## REVIEW

# The mechanism of sperm–egg interaction and the involvement of IZUMO1 in fusion

Naokazu Inoue, Masahito Ikawa and Masaru Okabe

An average human ejaculate contains over 100 million sperm, but only a few succeed in accomplishing the journey to an egg by migration through the female reproductive tract. Among these few sperm, only one participates in fertilization. There might be an ingenious molecular mechanism to ensure that the very best sperm fertilize an egg. However, recent gene disruption experiments in mice have revealed that many factors previously described as important for fertilization are largely dispensable. One could argue that the fertilization mechanism is made robust against gene disruptions. However, this is not likely, as there are already six different gene-disrupted mouse lines (*Calmegin*, *Adam1a*, *Adam2*, *Adam3*, *Ace* and *Pgap1*), all of which result in male sterility. The sperm from these animals are known to have defective zona-binding ability and at the same time lose oviduct-migrating ability. Concerning sperm–zona binding, the widely accepted involvement of sugar moiety on zona pellucida 3 (ZP3) is indicated to be dispensable by gene disruption experiments. Thus, the landscape of the mechanism of fertilization is revolving considerably. In the sperm–egg fusion process, CD9 on egg and IZUMO1 on sperm have emerged as essential factors. This review focuses on the mechanism of fertilization elucidated by gene-manipulated animals.

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**Keywords:** egg; fusion; gene-manipulation; interaction; IZUMO1; sperm; zona pellucida

## INTRODUCTION

Fertilization is the phenomenon in which sperm and egg find each other, interact and fuse. This phenomenon is essential for all plants and animals which engage in sexual reproduction. Many experiments have been performed, and papers published, using a vast number of phyla. Despite the biological importance of fertilization, the molecular mechanism of fertilization remains unknown. For example, in mammals, almost all the proteins reported to be important in sperm–egg interaction and fusion turned out to be not essentially required in fertilization under the light of gene disruption experiments<sup>1–5</sup> (Figure 1). The gene knockout experiments have negated the old scheme of fertilization mechanism, but at the same time, this technique has serendipitously made us aware of the existence of essential genes.

A tetraspanin family CD9 which is expressed ubiquitously in the body was disrupted to find the effect on immune systems. The disruption was performed in three different laboratories at the same time and all three laboratories found that both the *Cd9*-disrupted male and the female mice were healthy. However, surprisingly, the *Cd9*-null females were sterile. Somehow, the ubiquitously-expressed CD9 was not essential in almost all the body parts, but was essential for egg to fuse with sperm.<sup>6–8</sup> The role of CD9 in fertilization might not have been found for many years if the gene disruption technique not been applied.

This review focuses on the studies on fertilization using gene-manipulated animals and describes the history of various candidates before and after the gene disruption experiments. Due to limitations

on length, it was not possible to mention all the factors examined by gene-disrupted mouse lines. In this review, we aimed to provide a brief outline of the disappearance of the old scheme and an introduction to the totally new scheme which is emerging.

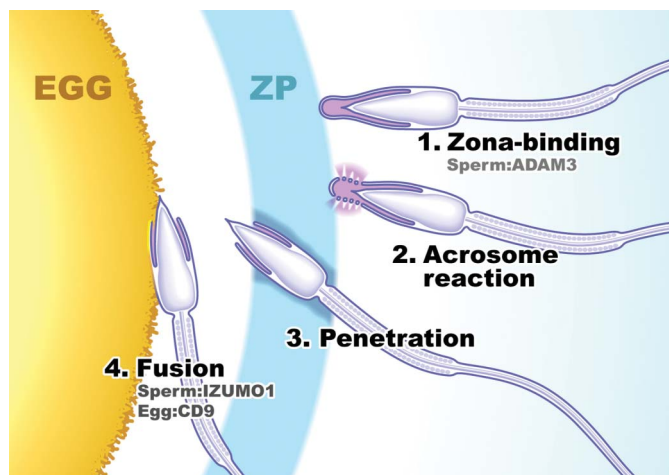
## CANDIDATE PROTEINS FOR SPERM–EGG INTERACTION

### Sperm–zona binding

There have been many papers indicating the involvement of acrosin on sperm-penetrating zona pellucida (ZP). Surprisingly, however, when *Acrosin*-knockout mice were produced, the sperm from the mutant mice fertilized eggs without difficulty.<sup>1</sup> This was just a prelude; all the so-called ‘important’ factors ( $\beta$ -1,4-galactosyltransferase, milk fat globule epidermal growth factor 8, zonadhesin, sperm adhesion molecule 1, etc.) have been demonstrated not to be essential in gene-disruption experiments.<sup>9–11</sup>

### Sperm–egg fusion

*Fertilin*. Until now, many monoclonal antibodies against sperm antigens were produced to elucidate the molecular mechanism of fertilization.<sup>12</sup> Among those, PH-30 was reported to inhibit sperm–egg fusion in *in vitro* fertilization in guinea pig. The antigen recognized by PH-30 was identified as fertilin that is composed of two glycosylated transmembrane subunits, fertilin- $\alpha$  (ADAM1B) and fertilin- $\beta$  (ADAM2). ADAM1B and ADAM2 make a heterodimer by non-covalent binding.<sup>13</sup> ADAM1B is conserved among broad species, albeit it is a pseudogene in human. ADAM is named after a disintegrin and metalloproteinase and forms family proteins consisting of multiple



**Figure 1** Mechanisms of sperm–egg interaction emerging from gene-manipulated animals. Some factors were found to be ‘essential’ after gene disruption. After being judged by gene disruption, ADAM3, IZUMO1 and CD9 are concluded to be indispensable factors in zona-binding on sperm, gamete fusion on sperm and on egg, respectively. As a whole, the explanation of sperm–egg interaction requires significant modification from the gene manipulation point of view. ZP consists of ZP1, ZP2 and ZP3 in mouse. Recently, it has been proved that sperm–egg recognition depends on the cleavage status of ZP2 by gene-manipulated mice. ADAM, a disintegrin and metalloprotease; ZP, zona pellucida.

domains: the prodomain, metalloprotease, disintegrin, cysteine-rich, epidermal growth factor-like, transmembrane and cytoplasmic tail domains. ADAM family proteins are fascinating proteins with important roles in cell adhesion, migration, proteolysis and signaling.<sup>14</sup> ADAM family proteins in human were reported up to number ADAM39 and are still expanding. In sperm from mouse testis, fertilin (ADAM1B/ADAM2) is distributed on the plasma membrane over the entire sperm head but is found only on the posterior head once sperm have passed through the epididymis. Moreover, during the transit from the testis to the epididymis, ADAM1B and ADAM2 are both proteolytically cleaved between the metalloprotease and disintegrin domains. Thus, in mature fertilization-competent sperm, the N-terminal of each fertilin subunit is the disintegrin domain.<sup>15</sup> Mouse ADAM2 has a peptide sequence similar to integrin-binding Arg–Gly–Asp domain. It was reported that when the recombinant protein of disintegrin domain added in *in vitro* fertilization assay, sperm–egg adhesion and fusion were inhibited.<sup>16</sup> Since the extracellular domain of ADAM1B contains a hydrophobic region that resembles the fusogenic region of viral fusion proteins, it was assumed that fertilin binds to an integrin ( $\alpha V\beta 3$  or  $\alpha 6\beta 1$  in mouse eggs) and thereby helps the sperm adhere to the surface of egg, which is a prerequisite for, and leads to, membrane fusion.<sup>17</sup> One of the ADAM proteins, meltrin- $\alpha$  (ADAM12), is reported to be involved in the formation of multinucleated myotubes.<sup>18</sup> These circumstantial data convinced many researchers to consider the fertilin as a genuine fusogenic factor in gamete fusion. If this is the case, the sperm without fertilin must fail to fuse with eggs. When fertilin was removed from sperm by eliminating one of the heterodimer genes *Adam2*, the male mice became infertile as expected, except that the disabled sperm function was largely a zona-binding ability rather than a fusing ability.<sup>19</sup> Moreover, the removal of fertilin by disrupting *Adam1b* gene, the sperm can fertilize eggs without fertilin.<sup>20</sup> Why do these two different fertilin knockout mouse lines show a completely different outcome? It is now understood that when the *Adam2* gene was deleted, a testicular type of fertilin (ADAM1A/ADAM2) was disrupted together with fertilin

(ADAM1B/ADAM2). Therefore, the apparent phenotype of ADAM2 was not directly derived from the disappearance of fertilin from sperm, but from the impaired formation of testicular type fertilin.<sup>19,20</sup>

Thus, the surprising outcome of gene disruption experiments is that fertilin is not essential for sperm–fertilizing ability despite considerable circumstantial evidence indicating that fertilin is the fusion protein.

*Cyritestin (ADAM3)*. ADAM3 is a 110-kDa protein in testis but is found to be a 42-kDa protein in epididymis similar to the case of ADAM1B and ADAM2.<sup>21</sup> An eight-residue peptide from the ADAM3 disintegrin loop sequence inhibits sperm–egg adhesion and fusion (80 and 90% of inhibition in adhesion and fusion, respectively). Therefore, ADAM3 was thought to be implicated in sperm–egg binding and fusion.<sup>22</sup> *Adam3*<sup>-/-</sup> sperm are drastically deficient in adhesion to the egg ZP (0.3% of wild-type) and to the egg plasma membrane (9% of wild-type) in *in vitro* fertilization assays. However, the gene disruption experiments indicated that ADAM3 is not essential for fusion (fertilization index remained the same as in the wild-type).<sup>23,24</sup>

*CD46*. Human CD46 is a ubiquitously-expressed protein known to protect cells from complement attack. Anderson *et al.* provide evidence that regulated generation of complement C3 fragments by acrosomal enzymes and the binding of these fragments by CD46 on sperm and complement receptor 1 on eggs may be an initial step in gamete interaction, leading to membrane fusion.<sup>25</sup> In fact, several anti-human CD46 monoclonal antibodies effectively inhibit fertilization in *in vitro* fertilization.<sup>26</sup> Interestingly, mouse CD46 was found only in testis and the protein was found on the inner acrosomal membrane of sperm.<sup>27</sup> The fact that CD46 is expressed only in testis in various mammalian species indicates the importance of CD46 in reproduction and encouraged us to make a *Cd46*-disrupted mouse line. Therefore, we produced mice carrying a null mutation in the *Cd46* gene. We found no difference in the fertilizing ability of sperm from *Cd46*-null mice in *in vivo* and *in vitro* systems. The only difference we discovered was the increase in spontaneous acrosome reaction in sperm from *Cd46*-null mice compared to that of the wild-type sperm. Differing from our expectation, without *Cd46*, the mice were healthy and were fertile in both sexes.<sup>27</sup>

*Cysteine-rich secretory protein (CRISP)*. CRISPs have molecular weights of about 20–30 kDa and are characterized by the presence of 16 conserved cysteine residues, 10 of which are clustered in the C-terminal domain of the molecule. CRISP1 and CRISP4 are enriched in the epididymis, CRISP2 is exclusively in developing spermatids in the testes, and CRISP3 is in wider tissue distribution than the other CRISPs.<sup>28,29</sup> When recombinant mouse CRISP1 and CRISP2 were added to eggs, they specifically bound to the fusogenic area of mouse eggs. The antibodies against these proteins significantly inhibit the fertilization ratio.<sup>30,31</sup> *Crisp1*-disrupted mouse line was produced and it was found that male and female *Crisp1*-null mice exhibited no differences in fertilizing ability compared to wild-type animals *in vivo*, even though the sperm from *Crisp1*-disrupted mice underwent lower tyrosine phosphorylation of capacitation and reduced ability to fertilize with egg *in vitro*.<sup>32</sup> It is clear that CRISP1 is dispensable for fertilization;<sup>32</sup> however, as for the other CRISP proteins, we must wait for the production of gene-disrupted mouse lines.

*MN9 antigen (equatorin)*. MN9 antigen/equatorin is an acrosomal protein found in various mammalian sperm, including human. It is

reported that equatorin translocates to the plasma membrane covering the equatorial region during the acrosome reaction. Equatorin on plasma membrane is considered to function in sperm–egg fusion, because MN9 antibody inhibits both *in vitro*<sup>33</sup> and *in vivo* fertilization systems<sup>34</sup> without affecting zona-penetrating ability of sperm. The *equatorin* gene is cloned<sup>35</sup> and has been identified as a type 1 transmembrane protein of a 40–60 kDa and *N,O*-sialoglycoprotein.

The production of an *equatorin*-deficient mouse line is underway (Toshimori K, 2010, per. commun.).

## ESSENTIAL PROTEINS FOR SPERM–EGG INTERACTION

### Sperm–zona binding

**ADAM3.** The first gene-disrupted mouse line that showed male infertility in spite of having a normal looking sperm is the case of *Calmegein* disruption reported from our laboratory.<sup>36</sup> Subsequently, five more genes were reported to cause male infertility (*Adam1a*, *Adam2*, *Adam3*, *Ace* and *Pgap1*). Interestingly, sperm from all six different gene-disrupted mouse lines are reported to have defects in zona-binding ability and oviduct-migrating ability.<sup>19,23,24,36–39</sup> This strongly indicates the involvement of a common factor in sperm migration into oviduct and zona-binding ability. As far as examined, all of these gene knockout mouse lines are reported to have defects in presenting ADAM3 on their surface except *Pgap1*. In *Pgap1* knockout mouse lines, the amount of ADAM3 on sperm was shown to be normal, but there is a chance that microdistribution of ADAM3 is affected by the elimination of PGAP1. Or else, the result of *Pgap1* disruption indicates that there is an essential factor for sperm–zona binding other than ADAM3. If not ADAM3, it is highly possible that the same factor is functioning in sperm migration into oviduct and in zona-binding ability because we could observe the dual phenotype in six different gene knockouts. The reason why so many gene disruptions result in the misplacement of ADAM3 on sperm is not clear. This is certainly an interesting question to investigate.

**ZP proteins.** ZP consists of ZP1, ZP2 and ZP3 in mouse (ZP4 is also found in human ZP). When sperm are mixed with eggs *in vitro*, many sperm are observed to bind to the zona. However, sperm ability to bind to zona is not necessary for sperm to fertilize eggs. It is reported that the *Adam1a*-disrupted mouse sperm show an impaired zona-binding ability but when the sperm are applied to the *in vitro* fertilization system using eggs with cumulus cells, the sperm with impaired zona-binding ability can fertilize eggs without any problem.<sup>38</sup> Ironically, although the zona-binding ability turned out to be dispensable, the mechanism of sperm binding to zona has been studied by many researchers. Mouse ZP3 has three important *O*-glycosylation sites, Ser-332, and Ser-334, and the mutation into these sites was predicted to abolish the sperm-binding ability by an *in vitro* translation system using somatic cells.<sup>40</sup> However, when the same experiment was performed *in vivo* using gene disruption and transgenic rescue with glycosylation incompetent mutant ZP3, the glycosylation in these sites were indicated to be not essential. It was also clarified that sperm-binding ability is largely regulated by the cleavage of ZP2 after fertilization.<sup>5</sup> However, as indicated above, the zona-binding ability of sperm seems to be not a critical characteristic for sperm to fertilize eggs.<sup>38</sup>

### Sperm–egg fusion

**CD9.** CD9 is a ubiquitously-expressed protein and is expected to function in various parts of the body. In 2000, three independent laboratories serendipitously and simultaneously found that a tetraspanin

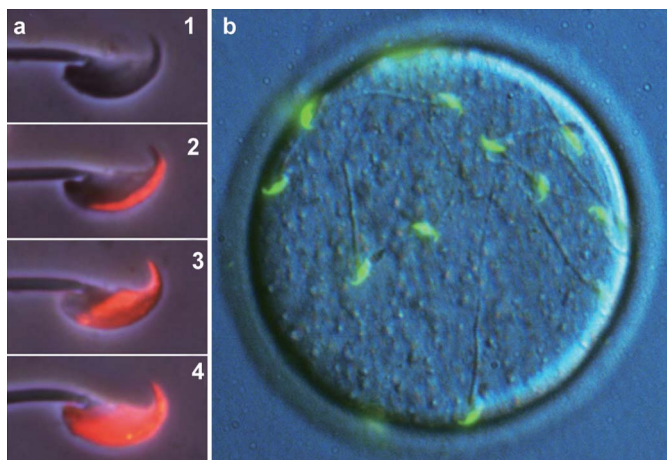
family CD9 on eggs was essential for eggs to fuse with sperm.<sup>6–8</sup> In *in vitro* fertilization assays, *Cd9*<sup>−/−</sup> eggs failed to fuse with sperm. Therefore, the eggs remain unfertilized and many sperm continuously penetrate the zona and result in the accumulation of unfused sperm in the perivitelline space. This defect is limited to the fusion process because the infertility of *Cd9*<sup>−/−</sup> eggs can be rescued by intracytoplasmic sperm injection.<sup>6</sup> Runge *et al.*<sup>41</sup> found that *Cd9*<sup>−/−</sup> eggs have an altered length, thickness and density of their microvilli by electron microscopy, suggesting that microvilli may participate in sperm–egg fusion. CD9 was the first fusion-related protein proved to be essential in the gene-manipulated animals.

**Glycosylphosphatidylinositol (GPI)-anchored protein.** Lipid microdomains called rafts are considered to provide specific fields on cells to facilitate many cellular processes such as signal transduction, membrane trafficking, cytoskeleton organization and pathogen entry. GPI-anchored proteins are one of the factors enriched in rafts. Coonrod *et al.* reported that treatment of eggs with PI-PLC significantly reduced sperm–egg binding and fusion.<sup>42</sup> When the GPI-anchoring process was demolished by egg-specific disruption of *Pig-a* gene using ZP3 promoter-driven *Cre* and loxP system, the female mice became infertile due to severely impaired fusing ability with sperm.<sup>43</sup> There are some indications of association of CD9 with GPI-anchored proteins,<sup>44</sup> but there is another report that tetraspanin web is distinct from raft microdomains.<sup>45</sup> Since both CD9 and GPI-anchored proteins are found on the egg surface, it would be interesting to know if there is an associated function between the CD9 and GPI-anchored proteins for fusion.

**IZUMO1.** We produced anti-mouse sperm monoclonal antibody OBF13 that inhibits the fusion process both *in vitro* and *in vivo*.<sup>46,47</sup> The antigen recognized by OBF13 was not identified for many years. However, it was recently identified by two-dimensional gel electrophoresis and subsequent immunoblotting and liquid chromatography-tandem mass spectrometry analysis. We named the antigen ‘Izumo’ after a Japanese shrine dedicated to marriage. The gene encodes a novel immunoglobulin superfamily type I membrane protein with one extracellular Ig domain. Recently, according to Ellerman *et al.*,<sup>48</sup> IZUMO proteins consist of four family proteins (IZUMO1 to IZUMO4). The N-terminal domain between signal peptides and Ig domain showed a significant homology to each other and was termed ‘Izumo domain’.

Acrosome-reacted sperm can be classified into three major groups by their IZUMO1-staining pattern: acrosomal cap, equatorial and whole-head (Figure 2, a2–a4). This indicates that during the acrosome reaction, IZUMO1 relocates from the anterior head of the sperm to the site(s) where fusion will take place. Since it is said that sperm launch fusion to egg at the equatorial segment, either equatorial or whole-head type IZUMO1 (Figure 2, a3 or a4) can contribute to sperm–egg fusion.

However, the question of whether or not IZUMO1 functions in fertilization could not be answered conclusive until the *Izumo1*-deficient mice were generated by homologous recombination. After producing *Izumo1*<sup>−/−</sup> mutant mice, we found they were healthy and showed no overt developmental abnormalities. As we expected, *Izumo1*<sup>−/−</sup> males became sterile despite normal mating behavior and ejaculated to form normal vaginal plugs. Moreover, the sperm penetrated the ZP without any problem but failed to fuse with eggs. This caused an accumulation of sperm in the perivitelline space of the eggs<sup>49</sup> (Figure 2b).



**Figure 2** Gamete fusion-related factor IZUMO1. **(a)** IZUMO1 is an acrosomal membrane protein that is not exposed before the completion of an acrosome reaction (1). Acrosome-reacted sperm can be classified into three major groups by their IZUMO1-staining pattern: acrosomal cap (2), equatorial (3) and whole-head (4). **(b)** Accumulation of many sperm in the perivitelline space of the eggs recovered from the females mated with *Izumo1*<sup>-/-</sup> males. Sperm in the perivitelline space were labeled with acrosome-reacted sperm-specific monoclonal antibody MN9.

Why did *Izumo1*<sup>-/-</sup> sperm fail to fuse with eggs? The acrosome reaction is a prerequisite for sperm to fuse with eggs. To help answer this, we examined the acrosomal status of *Izumo1*<sup>-/-</sup> sperm. In order to verify the acrosome reaction, we stained the sperm with MN9 monoclonal antibody which immunostains only to the equatorial segment of ‘acrosome-reacted’ sperm<sup>33</sup> (Figure 2b). As shown in the Figure 2b, the *Izumo1*<sup>-/-</sup> sperm was clearly stained with MN9. This indicated the *Izumo1*<sup>-/-</sup> sperm had undergone the acrosome reaction but failed to fuse with eggs.

We further examined whether the defect of *Izumo1*<sup>-/-</sup> sperm is limited to their fusing ability with eggs or whether it extends to later developmental stages. To address this question, we injected *Izumo1*<sup>-/-</sup> sperm directly into the cytoplasm of wild-type eggs and observed the ability of later development. Eggs injected with *Izumo1*<sup>-/-</sup> sperm were successfully activated and the eggs implanted normally. The embryos developed to term in a normal ratio.<sup>49</sup>

## CHARACTERIZATION OF IZUMO1 PROTEIN

### The structure of IZUMO1

As mentioned above, IZUMO1 has an Ig domain with 145 residues of N-terminal IZUMO domain. Electrophoresis under mildly denaturing conditions, followed by western blot analysis, showed that IZUMO1 and IZUMO3 formed complexes with other protein(s) on sperm. Studies using recombinant IZUMO1 constructs suggested the IZUMO1 domain possesses the ability to form dimers. IZUMO1 might be involved in organizing or stabilizing a multiprotein complex essential for the function of the membrane fusion machinery.<sup>48</sup>

IZUMO1 possess a well-conserved *N*-glycosylation site in the middle of an Ig loop among species. This site must be actually glycosylated because if we incubated mouse IZUMO1 from sperm with *N*-glycosidase, the molecular weight of IZUMO1 decreased to 50 kDa from its original 56 kDa. Since glycan composition is known to be involved in many molecular interaction mechanisms,<sup>50</sup> we tried to examine the role of *N*-glycan on IZUMO1. In order to answer this question, we produced mouse lines expressing mutated IZUMO1 in which the one hundred and eighty-third putative *N*-glycosylation site asparagine was

substituted to glutamine by site-directed mutagenesis under the testis-specific *Calmegein* promoter with rabbit  $\beta$ -globin polyadenylation signal. After we established N183Q-IZUMO1 males, we crossed these transgenic mouse lines with *Izumo1*<sup>-/-</sup> mice and produced a mouse lines which have sperm with no *N*-glycosylation site in IZUMO1. Although the litter sizes were smaller compared to the wild-type IZUMO1, the N183Q-IZUMO1 rescued the infertile phenotype back to a fertile one. The efficiency was low, but sperm from N183Q-IZUMO1 could fuse with eggs (Figure 3a). We extracted proteins from testis and sperm from the N183Q-IZUMO1 male mice and analyzed IZUMO1 by western blot analysis. The N183Q-IZUMO1 from testes was migrated to a 50-kDa band area due to the lack of *N*-linked glycan. However, N183Q-IZUMO1 from sperm, a severe fragmentation was observed which is not observed in wild-type IZUMO1. The major fragmented bands were observed in the ~30- and ~35-kDa areas (Figure 3b). Although N183Q-IZUMO1 could rescue the infertile phenotype, the amount of intact N183Q-IZUMO1 present on sperm was significantly small compared to that of wild-type IZUMO1 in spite of an abundance of N183Q-IZUMO1 in testis.<sup>51</sup> This indicates that glycosylation is not essential for the function of IZUMO1, but has a role in protecting it from fragmentation in cauda epididymis.

### IZUMO1-interacting protein

Since IZUMO1 has no ‘fusogenic’ peptide or ‘SNARE’-like structure in it, we considered the possibility that IZUMO1 might be one of the components forming fusogenic machinery on sperm. In order to search for IZUMO1-interacting proteins, we made a transgenic mouse line producing IZUMO1-His on sperm and introduced it to an *Izumo1*<sup>-/-</sup> background. This allowed us to immunoprecipitate IZUMO1 using anti-His antibody. The IZUMO1-interacting protein was purified from acrosome-intact sperm lysate using anti-His microbeads. We could find a specific 80-kDa band by silver staining in the purified fraction. After liquid chromatography–tandem mass spectrometry analysis, the protein was identified as ACE3 (angiotensin I-converting enzyme 3) (Figure 4a).<sup>52,53</sup>

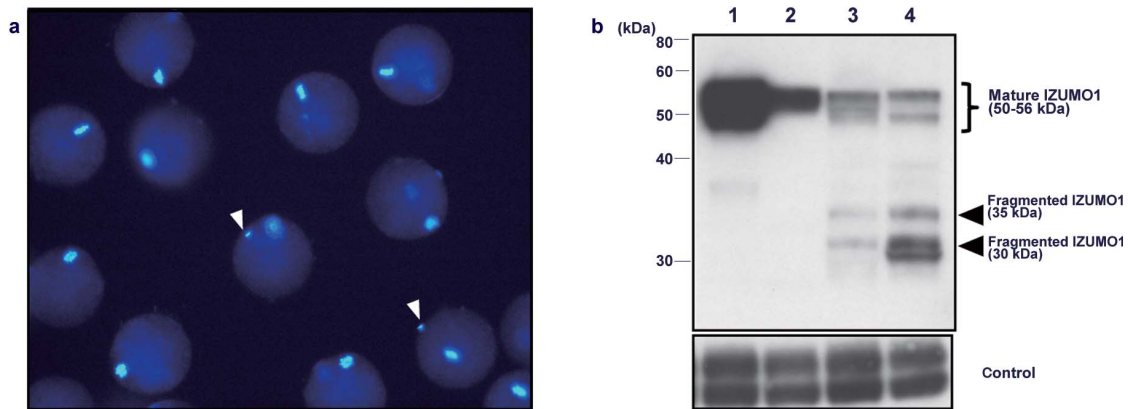
The immunofluorescent staining revealed that the ACE3 in fresh sperm localizes in the acrosomal cap area similar to SP56 and ACROSIN.<sup>54,55</sup> Although the ACE3 was detected in the acrosomal cap area colocalizing with IZUMO1 before acrosome reaction, it disappeared from acrosome-reacted sperm while IZUMO1 remained (Figure 4b).

As mentioned earlier, the real function of proteins on sperm cannot be judged unless the corresponding gene is genetically modified in the mouse. We therefore generated *Ace3*-deficient mice by homologous recombination. Differing from our expectation, the *Ace3*<sup>-/-</sup> mice showed signs of infertility both in males and in females. We analyzed the fertilizing ability of *Ace3*<sup>-/-</sup> sperm in the *in vitro* fertilization systems. Again, the *Ace3*<sup>-/-</sup> sperm showed normal fertilizing ability in our *in vitro* fertilization system using both cumulus-intact and cumulus-free eggs. These results suggest that ACE3 do bind to IZUMO1, but this characteristic nature is not required for sperm to fertilize eggs.<sup>53</sup>

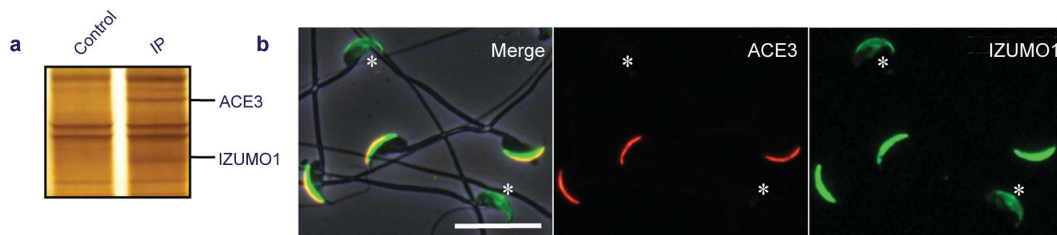
## FACTORS THAT AFFECT THE LOCALIZATION OF IZUMO1

### Sperm equatorial segment protein 1 (SPESP1)

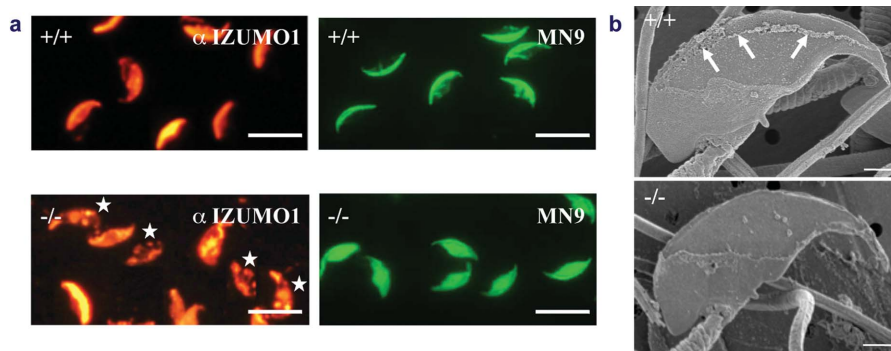
It is well accepted that sperm–egg fusion starts at the equatorial segment of sperm head. IZUMO1 can spread over the entire head covering the equatorial segment, but this does not explain the equatorial segment-restricted fusion.<sup>49</sup> There are various sperm proteins known to be distributed only in the equatorial segment. There are chances that



**Figure 3** N-linked glycan of IZUMO1 is not essential for fusion. **(a)** Comparison of the fusing ability of wild-type and N183Q-IZUMO1 sperm. The arrowheads indicate fused sperm. N183Q-IZUMO1 sperm are able to fuse with eggs, albeit in low yield (fusion index: 0.05 fused sperm/egg). **(b)** Fragmentation of N183Q-IZUMO1 protein in cauda epididymal sperm. N183Q-IZUMO1 is fragmented by protease in cauda epididymal sperm (filled arrowheads). Lane 1, Wild-type; Lane 2, *Izumo1*<sup>-/-</sup> *Izumo1*-Tg; Lane 3, *Izumo1*<sup>-/-</sup> *Izumo1*N183Q-Tg#1; Lane 4, *Izumo1*<sup>-/-</sup> *Izumo1*N183Q-Tg#2.



**Figure 4** Identification of IZUMO1-interacting proteins. **(a)** The purified IZUMO1 protein complex was separated by SDS-PAGE and then silver stained. Two specific 80- and 56-kDa bands appeared corresponding to ACE3 and IZUMO1, respectively. **(b)** Subcellular localization of ACE3 protein in mature sperm was examined in incubated cauda epididymal sperm. They were stained with anti-ACE3 (red) and anti-IZUMO1 (green) antibodies. The anti-ACE3 antibody stained acrosome-intact sperm head, but did not react to acrosome-reacted sperm (asterisk). Scale bar=20  $\mu$ m. ACE3, angiotensin I-converting enzyme 3; PAGE, polyacrylamide gel electrophoresis.



**Figure 5** SPESP1. **(a)** Many of the *Spesp1*-disrupted sperm showed abnormal spreading of IZUMO1 after acrosome reaction (indicated by asterisks in the figure). The localization pattern of MN9 antigen was also significantly affected by the disruption of *Spesp1*. Scale bars=10  $\mu$ m. **(b)** Scanning electron microscopy of *Spesp1*-deficient and wild-type sperm. In wild-type sperm, the plasma membrane in the acrosome cap area disappeared down to the border of the equatorial segment after acrosome reaction. However, in *Spesp1*-deficient sperm, the plasma membrane disappeared from a wider area. As a result, the inner acrosomal membrane was exposed in most of the equatorial segment area (indicated by arrowheads). Scale bars=1  $\mu$ m. SPESP1, sperm equatorial segment protein 1.

one of these proteins is functioning in sperm-egg fusion.<sup>35,56-58</sup> SPESP1 was one of the candidates functioning for fusion, because anti-SPESP1 antibody was known to inhibit human sperm from fusing with hamster eggs as well as inhibiting sperm-egg fusion in mouse.<sup>59</sup>

A mouse line lacking *Spesp1* was produced by homologous recombination and the fertilizing ability of male mice was examined.<sup>60</sup> The decreased fertilizing ability was found not only in *Spesp1*<sup>-/-</sup> sperm but also in *Spesp1*<sup>+/-</sup> sperm. Despite the normal number of ejaculated

sperm and normal motility, *Spesp1*-disrupted mouse had a decreased number of pups compared to wild-type mice. When we examined the localization of IZUMO1 in *Spesp1*-disrupted sperm, we found IZUMO1 localized in a broader area in a more spotty manner. The aberrant distribution of membrane proteins was more prominent in the case of MN9 antigen. In wild-type sperm, MN9 antigen did not spread to the equatorial segment before acrosome reaction, but when *Spesp1* was disrupted, MN9 antigen was not restricted to the acroso-

mal cap area, but to the equatorial segment even before the acrosome reaction (Figure 5a). When we observed the sperm using a scanning electron microscope, we found a complete removal of the equatorial segment membrane from all of the *Spesp1*<sup>-/-</sup> sperm (Figure 5b). This could be an artifact during the preparation of the sperm sample, but it is certain that the equatorial segment membrane in *Spesp1*<sup>-/-</sup> sperm becomes very fragile compared to that of wild-type sperm and affects the distribution of IZUMO1.<sup>60</sup>

### TSSK6

TSSK6 is a member of the testis-specific serine kinase and is expressed postmeiotically in male germ cells. The *Tssk6*-null mice are infertile producing mostly morphologically deformed sperm. However, the morphology is not the only cause for infertility. Thirteen percent of the sperm from the mice showed a normal morphology, but failed to fertilize eggs *in vitro* and failed to fuse with zona-free eggs. Interestingly, in *Tssk6*<sup>-/-</sup> sperm, the relocalization of IZUMO1 after acrosome reaction is impaired. It is assumed that polymerization of actin is required for this relocalization, because an inhibitor for actin polymerization blocks the relocalization of IZUMO1.<sup>61</sup> However, in *Tssk6*<sup>-/-</sup> sperm, polymerized actins disappeared from the midpiece and the posterior head of sperm. Thus, TSSK6 may regulate the localization of IZUMO1 by regulating the polymerization of actin after acrosome reaction.<sup>61</sup>

### CONCLUSION

Recent observations of infertile phenotypes derived from various gene-disrupted mouse lines have rendered the old scheme for the mechanism of fertilization obsolete. Many essential genes were required to express ADAM3 properly on the sperm surface, as shown by gene disruption experiments.<sup>19,23,24,36–38</sup> This indicates that ADAM3 is the key molecule in fertilization, but this scheme is not applicable in human because *Adam3* is a pseudogene in human.<sup>62</sup> Thus, although a partial mechanism has been clarified, a new (or modified) outline for the mechanism of fertilization must await the results of further gene knockout experiments. Concerning fusion mechanism, it might be important to broaden our focus to wider areas of the cell–cell fusion process such as the formation of myotubes, placenta, multinucleated osteoclasts and macrophages, for example. In any case, the clarification of the molecular mechanism of fertilization will benefit clinical treatment of infertility and will underpin the potential development of novel contraceptive strategies in the future.

### COMPETING FINANCIAL INTERESTS

The authors declare no competing financial interests.

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