Chapter 34
Involvement of Carbohydrate Residues of the Zona Pellucida in In Vitro Sperm Recognition in Pigs and Cattle

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Abstract The zona pellucida (ZP), which surrounds the mammalian oocyte, plays roles in species-selective sperm–oocyte interactions. In pigs and cattle, the ZP consists of ZP2, ZP3, and ZP4. Nonreducing terminal β-galactosyl (Gal) residues of neutral N-linked carbohydrate chains of the ZP are necessary for porcine sperm–ZP binding, and nonreducing terminal α-mannosyl (Man) residues of high-mannose-type chains are necessary for bovine sperm–ZP binding. Acrosome-intact porcine sperm prefer β-Gal, whereas acrosome-intact bovine sperm prefer α-Man, as shown using glycolipid analogues. The major N-linked chains of recombinant porcine and bovine ZP glycoproteins expressed using the baculovirus-Sf9 cell expression system are pauci- and high-mannose-type chains that are different in structure from the major neutral N-linked chains of the native porcine ZP but similar to those of the native bovine ZP. Porcine and bovine ZP3/ZP4 complexes coexpressed in Sf9 cells bind to bovine sperm but not to porcine sperm. Hybrid complexes consisting of native porcine ZP4 and recombinant porcine ZP3 bind to porcine sperm, whereas complexes consisting of native porcine ZP3 and recombinant porcine ZP4 do not bind to porcine sperm. These data indicate that the sugar preference of sperm is consistent with the nonreducing terminal residues of N-linked chains of sperm binding-active ZP glycoproteins and suggest that, in the in vitro assay system, the nonreducing terminal sugar residues are essential for species-selective recognition of sperm in pigs and cattle.

Keywords Baculovirus-Sf9 cell • N-linked chains • Sperm ligand • Zona pellucida
34.1 Introduction

Mammalian oocytes are surrounded by a transparent envelope called the zona pellucida (ZP), which is involved in several critical aspects of fertilization, including species-selective recognition of sperm. In many mammalian species, there are four ZP glycoproteins (ZPGs) (ZP1, ZP2, ZP3, ZP4) (Stetson et al. 2012), although there are three ZPGs (ZP1, ZP2, ZP3) in mice. The porcine and bovine ZPs are composed of three ZPGs (ZP2, ZP3, ZP4). All ZPGs contain the ZP domain, which consists of about 260 amino acids and contains eight conserved Cys residues (Jovine et al. 2005).

Many studies have proposed that in mice the carbohydrate chains of ZP3 play an essential role in sperm recognition (the glycan model). However, a series of studies using transgenic mice do not support the glycan model (Hoodbhoy and Dean 2004). Instead, the supramolecular complex model, in which the supramolecular structure of the ZP polypeptides is necessary for sperm recognition in mice, was proposed based on studies using transgenic mice rescued by the human ZP2 gene (Rankin et al. 2003). Furthermore, the domain-specific model, in which both protein and carbohydrate moieties in the domain(s) of ZPGs are involved in sperm recognition, was recently proposed (Clark 2011). More recently still, it was shown using transgenic mice that sperm–ZP binding is not necessary for fertilization of oocytes surrounded by the cumulus oophorus (Tokuhiro et al. 2012). Whether this finding is applicable to mammals other than mice remains to be determined.

One of our research questions is whether carbohydrate chains are involved in sperm–ZP binding in pigs and cattle. Here, we discuss this subject mainly by reviewing our biochemical studies.

34.2 Polypeptides of Porcine and Bovine ZPGs

The porcine ZP is approximately 16 μm in width and contains 30–33 ng glycoproteins. The bovine ZP is of similar size to the porcine ZP. The estimated ZP2/ZP3/ZP4 protein molar ratio is 1:6:6 in the porcine ZP and 1:2:1 in the bovine ZP. In mice, the estimated ZP1/ZP2/ZP3 molar ratio is 1:4:4, and ZP2 and ZP3 form a filamentous equimolar complex, whereas ZP1 crosslinks the ZP2/ZP3 complex (Greve and Wassarman 1985). Based on the similarity in the protein molar ratios, it appears that ZP architecture is similar in pigs and mice but bovine ZP architecture appears to differ from that of porcine and murine ZP.

Cys positions of the ZP domain are classified into two patterns: the ZP3 pattern and the ZP1/ZP2/ZP4 pattern (Fig. 34.1). Porcine ZP3 and ZP4 exhibit disulfide patterns different from the ZP3 and ZP1/ZP2/ZP4 patterns reported for mice, rats, humans, and fish (Kanai et al. 2008). The chick homologue of the mammalian ZP3 precursor protein has a pig-type ZP3 pattern (Han et al. 2010), whereas betaglycan has a pig-type ZP1/ZP2/ZP4 pattern (Lin et al. 2011), as revealed by X-ray crystallography. The two ZP3 disulfide bond patterns cause only subtle structural differences (Han et al. 2010). The subtle structural differences may affect the specificity of...
the interaction between ZP1/ZP2/ZP4 and ZP3 (Darie et al. 2004). When the disulfide bond pattern of ZP3 is of the pig type, the ZP1/ZP2/ZP4 pattern of the same species is also the pig type, and vice versa. The disulfide bond patterns might change during zona matrix formation, following processing at the consensus furin cleavage site.

### 34.3 N-Linked Carbohydrate Chains from Porcine and Bovine ZPGs

The N-linked chains of porcine ZP comprise neutral and acidic chains at a molar ratio of about 1:3. Both neutral chains and acidic chains comprise di-, tri-, and tetra-antennary complex-type chains with an α-fucosyl (Fuc) residue at the innermost N-acetylglucosamine (GlcNAc) (Nakano and Yonezawa 2001). A large proportion of neutral chains have nonreducing terminal β-galactosyl (Gal) residues.

In the bovine ZP, the major neutral N-linked chain consists of only one structure: the high-mannose-type chain containing five mannosyl (Man) residues (Man$_5$GlcNAc$_2$) (Katsumata et al. 1996). Sialylation is predominant in acidic chains of bovine ZP; in porcine ZP, sulfation of the N-acetyl lactosamine unit is more dominant.

### 34.4 Involvement of Carbohydrate Chains from Porcine and Bovine ZPGs in Sperm–ZP Binding

Neutral N-linked carbohydrate chains released from the porcine ZP3/ZP4 mixture retain their sperm-binding activities, with the tri- and tetra-antennary complex-type chains binding more strongly than the di-antennary complex-type chains...
In one study, the nonreducing terminal β-Gal residues of the complex-type N-linked chains were shown to be involved in sperm binding (Yonezawa et al. 2005a). Conversely, another study reported that the O-linked carbohydrate chains, and not the N-linked chains, released from the ZP3/ZP4 mixture inhibit sperm–ZP binding (Yurewicz et al. 1991). Therefore, both the N- and O-linked carbohydrate chains are thought to be sperm ligands. A polypeptide scaffold is necessary for the sperm-binding activity of carbohydrate chains, because the sperm-binding activities of carbohydrate chains released from polypeptides are much weaker than the activity of the ZP3/ZP4 mixture.

Porcine ZP3 and ZP4 have three N-linked chains, at Asn124, Asn146, and Asn271 in ZP3 and at Asn203, Asn220, and Asn333 in ZP4. Tri- and tetra-antennary chains are localized at Asn271 in ZP3 and at Asn220 in ZP4 (Nakano and Yonezawa 2001). ZP2 has five N-glycosylation sites: Asn84, Asn268, Asn316, Asn323, and Asn530 (von Witzendorff et al. 2005). A remarkable difference between the N-linked chain structures of ZP2 and ZP3/ZP4 is that ZP2 has a high-mannose-type chain containing five Man residues, probably located at Asn268.

Nonreducing terminal α-Man residues of the high-mannose-type chain play an essential role in bovine sperm–ZP binding (Amari et al. 2001). Sialic acid residues at the nonreducing ends of acidic N-linked or O-linked chains of bovine ZPGs are also involved in sperm binding (Velásquez et al. 2007). The N-glycosylation sites of bovine ZP3 and ZP4 have not yet been determined. The N-glycosylation sites of bovine ZP2 are Asn83, Asn191, and Asn527 (Ikeda et al. 2002).

Glycolipid analogue–possessing monosaccharides adsorbed on plastic wells show sperm-binding activity, but monosaccharide solution does not inhibit sperm–ZP binding (Takahashi et al. 2013). Two-dimensional coating of plastic wells with glycolipid analogue may cause the formation of a cluster of sugars, which is necessary for the high avidity for sperm binding. Acrosome-intact porcine sperm exhibit their strongest affinity for plastic wells coated with β-Gal and the second strongest affinity for β-N-acetylgalactosamine (GalNAc), whereas acrosome-intact bovine sperm exhibit their strongest binding affinities for α-Man and their second strongest affinity for β-glucose (Glc) and β-GlcNAc. These specificities are consistent with the nonreducing terminal sperm ligand sugar residues of porcine and bovine ZPs.

### 34.5 Sperm Binding-Active Regions of Porcine and Bovine ZPGs

We identified the sperm binding-active fragment of porcine ZP4 as an N-terminal region (Asp137 to Lys247) containing two N-linked chains (Nakano and Yonezawa 2001). Porcine ZP4 purified by reverse-phase HPLC is actually contaminated with ZP3, and formation of a heterocomplex of ZP3 and ZP4 is essential for the sperm-binding activity of the glycoproteins (Yurewicz et al. 1998). We established an expression system for recombinant porcine ZPGs (rpZPGs) using baculovirus-Sf9...
cells to obtain ZP4 without contamination of ZP3 (Yonezawa et al. 2005b). The major structures of the N-linked chains of rpZPGs are believed to be pauci- and high-mannose-type chains with or without a Fuc residue at the innermost GlcNAc. They have nonreducing terminal α-Man residues and are thus similar in structure to the sperm ligand N-linked chain of bovine ZPGs. The rpZP3/rpZP4 mixture coexpressed by Sf9 cells shows binding activity toward bovine sperm but not to porcine sperm (Fig. 34.2). The sperm-binding activity of rpZP4 is much weaker than that of rpZP3/rpZP4 (Yonezawa et al. 2005b). This study supports the previous report by Yurewicz et al. (1998).

To reexamine the sperm binding-active region of porcine ZP4, rpZP4 mutants with an Asp substitution at Asn203, Asn220, or Asn333 were coexpressed with rpZP3 in Sf9 cells (Yonezawa et al. 2005b). The mutation at Asn333 does not have an effect on the sperm-binding activity of the rpZP3/rpZP4 mixture, whereas the mutations at Asn203 and Asn220 reduce the sperm-binding activity secondarily and primarily among the three sites, respectively. These results are in agreement with our previous finding that the N-terminal fragment containing two N-linked chains at Asn203 and Asn220 exhibits sperm-binding activity, and that the tri- and tetra-antennary chains are localized at Asn220 in porcine ZP4.

Porcine sperm bind to native porcine ZPGs, but not to rpZPGs, probably because of differences in carbohydrate structures between native and recombinant ZPGs. A mixture of rpZP3 and native porcine ZP4 binds to the acrosomal region of porcine

![Fig. 34.2 Schematic representation of the specificity in recognition between sperm and zona pellucida (ZP) glycoproteins. Porcine sperm bind to the heterocomplex of porcine ZP3 and ZP4 in which ZP4 has complex-type N-linked chains (closed circles in green) but not to the heterocomplexes in which ZP4 has pauci- and high-mannose-type chains (closed diamonds in red). On the other hand, bovine sperm bind to the heterocomplexes in which ZP4 has pauci- and high-mannose-type chains.](image-url)
sperm and inhibits porcine sperm–ZP binding (Fig. 34.2) (Yonezawa et al. 2012). A mixture of native porcine ZP3 and rpZP4 does not inhibit the binding, although the mixture inhibits bovine sperm–ZP binding. This study indicated that native ZP4, but not rZP4, is necessary for the binding activity of the porcine ZP3/ZP4 complex toward porcine sperm and further suggested that the carbohydrate structures of ZP4 in the porcine ZP3/ZP4 complex are responsible for the porcine sperm-binding activity of the complex. Thus, in the present model for porcine sperm–ZP binding, the N-linked chains with nonreducing terminal \( \beta \)-Gal residues linked to the N-terminal region of ZP4 form a sperm-binding domain in the ZP3/ZP4 heterocomplex.

ZP4 has the highest sperm-binding activity among bovine ZPGs purified from ovaries, while ZP2 and ZP3 have weak but significant sperm-binding activities (Yonezawa et al. 2001). In the case of bovine ZPGs, ZP4 is not completely purified by reverse-phase HPLC and is contaminated with ZP3. To obtain ZP3 and ZP4 without contamination of other ZPGs, we established an expression system for bovine ZPGs using the baculovirus-Sf9 cell system (Kanai et al. 2007). Recombinant bovine ZP3 (rbZP3) and rbZP4 do not show sperm-binding activity, whereas the rbZP3/rbZP4 complex does show sperm-binding activity. The region from the N-terminus to the trefoil domain of rbZP4 is dispensable for formation of the complex with rbZP3 and for the sperm-binding activity of the complex. More detailed analysis of the sperm-binding region of bovine ZP3/ZP4 has not been accomplished.

### 34.6 Conclusion

Our studies indicate that porcine and bovine sperm proteins recognize \( \beta \)-Gal and \( \alpha \)-Man residues, respectively, at the nonreducing ends of carbohydrate chains linked to the ZP3/ZP4 complex. In pigs and cattle, ZP3 and ZP4 form a framework on which carbohydrate chains active in sperm binding can exhibit high avidity for sperm. To prove the significance of the carbohydrate chains in sperm binding, the sperm proteins that bind specifically to \( \beta \)-Gal in pigs and \( \alpha \)-Man in cattle need to be identified and characterized.

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