Localized Surface Antigens of Guinea Pig Sperm 
Migrate to New Regions Prior to Fertilization

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ABSTRACT We have previously defined distinct localizations of antigens on the surface of the guinea pig sperm using monoclonal antibodies. In the present study we have demonstrated that these antigen localizations are dynamic and can be altered during changes in the functional state of the sperm. Before the sperm is capable of fertilizing the egg, it must undergo capacitation and an exocytic event, the acrosome reaction. Prior to capacitation, the antigen recognized by the monoclonal antibody, PT-1, was restricted to the posterior tail region (principal piece and end piece). After incubation in capacitating media at 37°C for 1 h, 100% of the sperm population showed migration of the PT-1 antigen onto the anterior tail. This redistribution of surface antigen resulted from a migration of the surface molecules originally present on the posterior tail. It did not occur in the presence of metabolic poisons or when tail-beating was prevented. It was temperature-dependent, and did not require exogenous Ca²⁺. Since the PT-1 antigen is freely diffusing on the posterior tail before migration, the mechanism of redistribution could involve the alteration of a presumptive membrane barrier. In addition, we observed the redistribution of a second surface antigen after the acrosome reaction. The antigen recognized by the monoclonal antibody, PH-20, was localized exclusively in the posterior head region of acrosome-intact sperm. Within 7–10 min of induction of the acrosome reaction with Ca²⁺ and A23187, 90–100% of the acrosome-reacted sperm population no longer demonstrated binding of the PH-20 antibody on the posterior head, but showed binding instead on the inner acrosomal membrane. This redistribution of the PH-20 antigen also resulted from the migration of pre-existing surface molecules, but did not appear to require energy. The migration of PH-20 antigen was a selective process; other antigens localized to the posterior head region did not leave the posterior head after the acrosome reaction. These rearrangements of cell surface molecules may act to regulate cell surface function during fertilization.

The arrangement of molecules into discrete domains on the surface of cells is now known to be of widespread occurrence (1–11). Membrane topography may reflect the functional geometry of the cell; for example, in intestinal epithelial cells digestive enzymes are enriched on the apical surface (12). Alternatively, the topographical distribution of cell surface molecules may control their function by determining the environment of the molecule or interactions between surface molecules.

The sperm cell is one of the mammalian cell types in which surface molecules are localized in domains. Using monoclonal antibodies to the guinea pig sperm surface, we have demonstrated that there are a minimum of 12 different antigens localized in five different patterns (10). 11 of these antigens have been identified as proteins. Evidence from other investigators indicates that these surface domains also differ in their lipid composition (5).

In this paper, we describe two topographical rearrangements of localized cell surface molecules. The rearrangements occurred before fertilization during steps that altered the functional state of the sperm. When mammalian sperm are either ejaculated or removed from the cauda epididymis they are unable to fertilize eggs. In the female reproductive tract, sperm undergo a set of alterations, collectively called capacitation, that makes them competent for fertilization. This process of capacitation can be mimicked in vitro by incubation in defined media. Capacitation leads to the exocytotic acrosome reaction and a change in motility pattern of the sperm. Only
those sperm which have acrosome-reacted are able to fuse with eggs (13, 14). During capacitation and after the acrosome reaction of guinea pig sperm, we observed a change in localization patterns of two different surface antigens from the initial localized pattern to a new localized pattern.

Redistribution or rearrangements of surface topography can theoretically occur by a variety of mechanisms, including the local insertion and elimination of molecules or change in accessibility of binding sites to the external surface by masking or unmasking. Migration of molecules already present on the surface may also be responsible for reordering cell surface topography. Both of the rearrangements described here resulted from the migration of pre-existing surface molecules. We have investigated a set of parameters that characterize these migrations.

**MATERIALS AND METHODS**

**Materials:** Monoclonal antibodies (MAbs) were generated by a fusion between spleen cells from immunized C57 Bl/6 female mice and P3-NS1/Aig 4-1 myeloma cells as described (10, 15). Antibody subclasses were determined by Ouchterlony assays: PT-1 is an IgG2b and PH-20 belongs to the subclass IgG1. The second antibodies used were fluorescein isothiocyanate (FITC)-Fab' or FITC-Fab goat anti-mouse IgG from Cappel Laboratories, Cochranville, PA. Erythro-9-(2-hydroxy-3-nunyl)adenine (EHNA) was purchased from Buckberg Lab Animals, Tomkins Cove, NY.

**Media:** The capacitating medium used was minimum culture medium (MCM) that contained 102.3 mM NaCl, 25.1 mM NaHCO3, 0.25 mM sodium pyruvate, and 21.7 mM sodium lactate. The MCM also contained 1.7 mM CaCl2 (Ca++ MCM) or 1.0 mM MgCl2 (Mg++ MCM) (16). In some cases 1.0 mM CaCl2 or 1.0 mM MgCl2 was added to the MCM to increase sperm survival (17). Other media used were Mg++ HEPES (140 mM NaCl, 4 mM KCl, 4 mM HEPES, pH 7.4, 10 mM glucose, 2 mM MgCl2; and 0.1 mM EGTA) and Ca++ HEPES (Mg++ HEPES, with 2 mM CaCl2 instead of MgCl2 and EGTA) (18).

**Immunofluorescence:** All immunofluorescent staining was carried out on live sperm or sperm fixed in 1.5% formaldehyde (made fresh). Fixed cells were washed through 5 ml heat-inactivated horse serum. Incubations in each of the MCM-based capacitating media tested: Mg++/Ca++ MCM or Mg++ K+ MCM + 1.0 mM NaCl, 1.0 mM MgCl2, 1.0 mM EGTA, 0.25 mM sodium pyruvate, and 21.7 mM sodium lactate. The MCM also contained 1.7 mM CaCl2 (Ca++ MCM) or 1.0 mM MgCl2 (Mg++ MCM) (16). In some cases 1.0 mM CaCl2 or 1.0 mM MgCl2 was added to the MCM to increase sperm survival (17). Other media used were Mg++ HEPES (140 mM NaCl, 4 mM KCl, 4 mM HEPES, pH 7.4, 10 mM glucose, 2 mM MgCl2; and 0.1 mM EGTA) and Ca++ HEPES (Mg++ HEPES, with 2 mM CaCl2 instead of MgCl2 and EGTA) (18).

**Incubations:** To test for the migration of the PT-1 antigen, we incubated 5 × 104 sperm (unwashed or after a wash in 0.9% NaCl) in 0.5 ml of the appropriate media in a temperature-controlled water bath or in a Lab-Line Orbit Environ-Shaker 18 (Lab-Line Instruments, Inc., Melrose Park, IL), rotating at 60 rpm. To test for migration of the PH-20 antigen, we incubated sperm at concentrations ranging from 4 × 108 to 107/ml under conditions leading to the acrosome reaction (compare below).

To test whether the PT-1 or PH-20 MAbs dissociate and reassociate during the incubations, we incubated cells labeled with the MAb and a goat FAb-FITC anti-mouse IgG in the presence of excess MAb in the medium. The amounts of excess MAb added to the medium and the amounts of MAb bound to the cells were determined. The concentrations in culture supernatants containing MAbs were followed by washing through 1 ml of horse serum and incubation in an FITC-conjugated secondary antibody. Further details have been described elsewhere (15). Patterns were observed on live swimming sperm. For photography, live-stained sperm swimming was stopped by the addition of 1 µg/ml A23187 to freshly obtained sperm suspended in Ca++/Mg++ HEPS at 37°C. Sperm were scored as acrosome-intact or acrosome-reacted using phase-contrast optics.

**RESULTS**

Two different antigens that are localized in domains on the surface of guinea pig sperm migrate into new regions of the sperm surface during the steps that lead up to fertilization.

**Migration of the PT-1 Antigen**

The first example of a migrating antigen is a protein characterized as having a sedimentation coefficient of 6.8S on octyglylucoside-containing sucrose density gradients (20). The 6.8S protein is recognized by the monoclonal antibody PT-1 and is limited to the surface of the posterior tail (principle piece and end piece) of sperm freshly taken from the cauda epididymis or vas deferens (Fig. 1, A and B). The binding pattern of PT-1 MAb is the same whether sperm are stained live at room temperature using a monoclonal FITC-Fab goat anti-mouse IgG or if they are stained live at 4°C or after fixation in 1.5-4% formaldehyde (15). When guinea pig sperm from the cauda epididymis were incubated in MCM at 37°C, the pattern of PT-1 antibody binding was changed to include the anterior tail (midpiece) of the sperm surface. Migration of the PT-1 antigen onto the anterior tail occurred in each of the MCM-based capacitating media tested: Mg++ MCM, Ca++ MCM, or Mg++ MCM + 1.0 mM NaCl (Mg++ K+ MCM), and in Mg++ K+ MCM containing 1 mM EGTA. Progressive stages in antigen migration were observed. During early stages in migration, the fluorescence on the anterior tail appeared to be dimmer or equal in brightness to the posterior tail region (Fig. 1, C and D). After longer periods of incubation many of the sperm showed antibody accumulation in the anterior tail region (Fig. 1, E and F). The exact percentage of cells showing accumulation varied with the incubation conditions but sometimes included >98% of the sperm in a population. However, under no conditions tested, even after 8 h of incubation, did the antigen completely leave the posterior tail region, or migrate onto the surface of the head.

To show that this change in antibody binding pattern was the result of the posterior tail antigen moving, and not new insertion or unmasking of PT-1 binding epitopes already on the anterior tail, we tagged the antigen with the MAB before migration. In this experiment, sperm were allowed to bind the PT-1 antibody (at 24°C where no migration occurred). Unbound monovalent antibody was washed away, cells were incubated in MCM at 37°C for 2 h, fixed in formaldehyde and stained with an FITC-conjugated Fab fragment of goat anti-mouse IgG. 100% of the sperm stained in the anterior tail region, showing that the PT-1 antigen moves onto the anterior tail with the MAB attached. To test the possibility that the attached MAB dissociates from the posterior tail antigen and re-binds to new epitopes on the anterior tail surface, we performed another experiment. Sperm, labeled with the PT-1 MAB followed by FITC-Fab goat anti-mouse IgG, were incubated in MCM containing a 100-fold excess of free PT-1 MAB for 2 h at 37°C and were then fixed. If, during 1 Abbreviations used in this paper: CCP, carbonyl cyanide m-chlorophenylhydrazone; EHNA, erythro-9-(2-hydroxy-3-nonyl)adenine; FITC, fluorescein isothiocyanate; IAMA, inner acrosomal membrane; MAB, monoclonal antibody; MCM, minimum culture medium; PVP, polyvinylpyrrolidone.
the incubation in MCM, new molecules were inserted or previously masked epitopes on the anterior tail were revealed, they would bind the excess PT-1 MAb in the medium. Since this excess PT-1 had no fluorescent tag attached, no fluorescence would be seen on the anterior tail. Fluorescence, however, was observed on the anterior tail of all the sperm. On the basis of these experiments, we concluded that the PT-1 antigen actually migrates onto the anterior tail.

The PT-1 antigen migration was temperature-dependent. After 30 min of incubation at 37°C, over 50% of sperm...
showed binding of PT-1 to the anterior tail region (Fig. 2). After 1 h, 95% of the sperm demonstrated migration of the antigen onto the anterior tail region. In the same experiment, the timing of migration was slowed by lowering the incubation temperature to 33°C, and, at 25°C, no antigen migration was detected even after 2 h of incubation (Fig. 2).

PT-1 antigen migration is also an energy-requiring process. When cells were incubated at 37°C in MCM in the presence of metabolic poisons, carbonyl cyanide m-chlorophenylhydrazone (CCCP) or 2,4-dinitrophenol, the PT-1 antigen showed little or no migration into the anterior tail domain (Fig. 3). Since treatment with these compounds resulted in the cessation of sperm motility, it is possible that sperm motility itself is required for antigen migration. To test this possibility we used other conditions to prevent sperm motility. In the first set of experiments, we used EHNA to inhibit the dynein ATPase necessary for flagellar beating (21, 22). When EHNA was added to a suspension of guinea pig sperm, swimming was blocked. As long as EHNA was present, and swimming had stopped, the PT-1 antigen did not migrate onto the anterior tail. If EHNA was added at 30 min, when a proportion of the cells already showed antigen migration onto the anterior tail, migration of the antigen on the surface of the remainder of the cells was prevented (Fig. 4). The effect of EHNA was reversible. If sperm were washed out of EHNA they would resume swimming and the antigen would migrate onto the anterior tail.

Although EHNA is a useful inhibitor for preventing sperm motility, it may affect other cell enzymes. Therefore, we used an alternative method for immobilizing sperm, by suspending them in a medium of high enough viscosity to physically

Figure 2 Temperature-dependence of PT-1 antigen migration. Sperm from one animal were incubated in Mg²⁺-MCM + 1 mM KCl. Percentage of cells showing binding of PT-1 antibody to the anterior tail region after incubation at 37°C (△), 33°C (○), or 25°C (●). After incubation, cells were fixed and stained by indirect immunofluorescence and 50 sperm were scored for the region of fluorescence for each time point. All cells showing fluorescence on the anterior tail are scored as migrated.

Figure 3 Inhibition of PT-1 antigen migration by metabolic poisons. Percentage of cells showing binding of PT-1 antibody to the anterior tail region after incubation in Mg²⁺-MCM + 1 mM KCl in the presence of 1 mM CCCP (○), or 10 mM 2,4-dinitrophenol (DNP) (●), and in both cases 1% dimethyl sulfoxide. Migration was assayed as in Fig. 2. The (no additions) control containing 1% dimethylsulfoxide is an average of two different experiments. (△) Migration of the PT-1 antigen was not affected by the presence of 1% dimethylsulfoxide.

Figure 4 Inhibition of PT-1 antigen migration by EHNA. Percentage of cells showing binding of PT-1 MAb to the anterior tail during incubation in Mg²⁺-MCM + 1 mM KCl in the presence of 4 mM EHNA (△) compared to a control with no additions (△). EHNA is added to the incubation medium 30 min after putting the cells at 37°C. At 30 min, ~20% of the cells showed PT-1 antigen migration onto the anterior tail; after the addition of EHNA, that percentage remains the same. Migration was assayed as described in Fig. 2. In this experiment the migration of PT-1 antigen in the control was slower than usual (cf. Figs. 2, 3, and 5), representing an observed variability of the kinetics of migration in different sperm samples.
prevent tail-beating. Sperm suspended in 10% polyvinylpyrrolidone 360 (PVP-360) did not swim and the PT-1 antigen did not migrate. As with the EHNA, this effect was reversible; when cells were washed out of PVP-360, both sperm swimming and antigen migration resumed (Fig. 5). Sperm were held in the nonmotile condition in PVP-360 for at least 1.5 h without affecting the subsequent migration of the PT-1 antigen (Fig. 5).

Migration of the PH-20 Antigen

As shown in Fig. 7, the PH-20 monoclonal antibody binds to the posterior head surface of live sperm freshly removed from the cauda epididymis (Fig. 7, A and B). Immunoprecipitation with the PH-20 MAb from detergent extracts of 125I surface-labeled sperm yields three labeled bands with Mr 66,000, 48,000, and 41,000 (10). The migration of the PH-20 antigen immediately follows the acrosome reaction. During the acrosome reaction there is a fusion, shown in Fig. 6, of the outer region of acrosomal membrane with the overlying region of the plasma membrane. The most proximal fusion results in a confluency between the acrosomal membrane and the plasma membrane. Distal to this confluency, hybrid vesicles of acrosomal and plasma membrane are released from the sperm along with the acrosomal contents. Therefore acrosome-reacted sperm are surrounded by a cell membrane that is a hybrid between plasma membrane and the remaining acrosomal membrane, composed primarily of the inner region of the acrosomal membrane (Fig. 6).

After the acrosome reaction, the PH-20 antigen migrated from the posterior head region onto the inner acrosomal membrane (IAM) (Fig. 7, A, B, E, and F). The migration to the IAM occurred regardless of whether the acrosome reaction was induced by capacitation in MCM or by the addition of A23187 to Ca2+ HEPES medium without prior capacitation. The kinetics of migration were followed after induction of the acrosome reaction with A23187. In the results shown in Fig. 8, 98% of the sperm were acrosome-reacted within 7 min of the addition of A23187; by 10 min, 100% of the acrosome-reacted sperm showed binding of PH-20 MAb bound only to the IAM. In repeated experiments with this protocol, the kinetics sometimes showed that as early as 7 min, 100% of the acrosome-reacted sperm had PH-20 MAb bound only to the IAM. During the migration, intermediate stages can be detected on the cells, with antigen present on both the posterior head and IAM, as shown in Fig. 8.

The same two experiments, establishing that the PT-1 antigen actually migrates, were done to show that the PH-20 antigen also migrates. If PH-20 MAb was bound to sperm before the acrosome reaction, the PH-20 antigen moved, with the MAb attached, to the IAM on all the cells. If sperm, with both PH-20 MAb and FITC-Fab goat anti-mouse Igs bound to them, were incubated in medium that contained a 50-fold excess of free PH-20 MAb, and then allowed to acrosome react, fluorescence was seen on the IAM of 100% of the sperm. Thus, we concluded that the PH-20 antigen migrates to the IAM.

Other antigens present on the surface of the posterior head did not migrate to the IAM after the acrosome reaction. Antigens that remained on the posterior head of acrosome-reacted sperm included posterior head antigens recognized by the monoclonal antibodies PH-1, PH-10, and PH-30. The binding pattern of PH-10 is shown before (Fig. 7, C and D) and after (Fig. 7, G and H) the acrosome reaction.

Unlike the migration of the PT-1 antigen, PH-20 antigen

![Figure 5](https://jcb.rupress.org/figure/5)

**Figure 5** Percentage of cells showing PT-1 MAb binding to the anterior tail region in the absence (Δ) or presence of 10% PVP (○) and after sperm have been washed out of PVP (●). In the presence of PVP no antibody binding is observed in the anterior tail region. When PVP is washed out, migration of the PT-1 antigen resumes. The times at which migration was scored after PVP removal are shown in parentheses.

![Figure 6](https://jcb.rupress.org/figure/6)

**Figure 6** A diagram showing what happens to the membranes of the mammalian sperm during the acrosome reaction. The plasma membrane of the anterior head fuses with the underlying acrosomal membrane except in a thin strip around the equator of the head (the equatorial region). Hybrid plasma-acrosomal membrane vesicles are released leaving the posterior head plasma membrane in continuity with the inner acrosomal membrane through the equatorial region.

![Diagram](https://jcb.rupress.org/figure/diagram)

**Diagram** A diagram showing what happens to the membranes of the mammalian sperm during the acrosome reaction. The plasma membrane of the anterior head fuses with the underlying acrosomal membrane except in a thin strip around the equator of the head (the equatorial region). Hybrid plasma-acrosomal membrane vesicles are released leaving the posterior head plasma membrane in continuity with the inner acrosomal membrane through the equatorial region.
FIGURE 7 Antibody binding patterns of the PH-20 and PH-10 MAbs on acrosome-intact and acrosome-reacted cells. (A) PH-20 indirect immunofluorescence and (B) phase contrast of acrosome-intact sperm, live staining. Bar, 5 μm. × 1,600. (C and D) PH-10 immunofluorescence on acrosome-intact sperm, live staining. Both antibodies bind exclusively to the posterior head. Bar, 5 μm. × 1,600. (E and F) PH-20 immunofluorescence on acrosome-reacted sperm, capacitated in Ca²⁺-MCM, live staining. Bar, 5 μm. × 1,700. (G and H) PH-10 immunofluorescence on acrosome-reacted sperm, capacitated in Ca²⁺-MCM, live staining. Bar, 5 μm. × 1,600.

FIGURE 8 The kinetics of migration of the PH-20 antigen on cells undergoing the acrosome reaction after addition of A23187. Percentage of acrosome-reacted cells (Δ); percentage of acrosome-reacted cells that show binding of PH-20 antibody to the inner acrosomal membrane but not to the posterior head (○); percentage of acrosome-reacted cells that show binding of PH-20 antibody to both the inner acrosomal membrane and the posterior head (□). Sperm, labeled with the MAb PH-20, are resuspended in Ca²⁺-HEPES at 37°C and 1 μg/ml A23187 is added at zero time. Aliquots are fixed at the times shown and examined for acrosome reaction and fluorescence. 50–100 cells are scored for each point. At zero time there is a background level of 4% acrosome-reacted sperm. By 3 min following the addition of ionophore, 94% of the sperm are acrosome-reacted, and within 7 min 84% of the acrosome-reacted sperm demonstrate binding of PH-20 on the IAM, but not in the posterior head region.

Migration did not appear to require energy. When ATP synthesis was blocked, the acrosome reaction and migration of the PH-20 antigen still occurred. If sperm were incubated in CCCP and 2-deoxyglucose, alone or in combination, sperm motility was completely inhibited. The addition of A23187 in the presence of these drugs induced the acrosome reaction and the subsequent migration of the PH-20 antigen onto the IAM (Table I).

Divalent second antibody was tested after the binding of PH-20 MAb to see if it could prevent the migration of the PH-20 antigen. We expected the divalent second antibody, a polyclonal FITC-F(ab')₂ anti-mouse IgG, to cross-link the antigen-monoclonal antibody complexes on the posterior head surface of sperm, and perhaps build up a network that would be unable to migrate after the acrosome reaction. When sperm with PH-20 monoclonal antibody and F(ab')₂ goat anti-mouse bound to them were induced to acrosome react, the PH-20 antigen remained in the posterior head and did not migrate to the IAM (Table I).

DISCUSSION
Membranes are a dynamic assembly of proteins and lipids that are able to respond to developmental and environmental signals to modify specific cellular functions. One response that they exhibit is a change of membrane topography or the redistribution of cell surface molecules into or out of specific domains. In some cases, surface molecules are redistributed by the actual migration of surface antigens in the plane of the membrane. Lateral migrations may be stimulated by ligand binding or occur in the absence of ligands. Ligand-stimulated migrations are evident in capping phenomena (23) and also include the movement of concanavalin A receptors to the...
cleavage furrow of cells (24) and the movement of several
tail-beating was prevented by incubation with the drug EHNA
or by keeping the sperm in a highly viscous solution of PVP-
360. Prevention of antigen migration by preventing tail-beat-
ing could indicate that tail-beating is somehow involved in
providing a force for PT-1 antigen relocation. Alternatively,
tail-beating may in some unforeseen way alter a presumptive
membrane barrier or maintain some cellular metabolic bal-
ance that allows migration to occur.

In contrast to PT-1 antigen migration, PH-20 antigen mi-
gation will occur in the presence of CCCP and 2-deoxyglu-
cose to block ATP synthesis. We cannot exclude the possibility
that residual ATP is used for the migration. However, as with
the relocation of lectin receptors in regions of cell-cell contact
(2), the migration of PH-20 antigen onto the IAM does not
appear to require cellular energy. Redistribution of the PH-
20 antigen in the presence of metabolic inhibitors is consistent
with models involving diffusion and trapping (2, 4) or ther-
modynamic partitioning.

The movement of PH-20 antigen did not result from a
mass flow of posterior head membrane components since it
is a selective process. Only the PH-20 antigen was shown to
move onto the IAM; other antigens remained behind in the
posterior head region.

The migrations of the surface antigens PT-1 and PH-20 are
each coincident with a change in the functional state of the
sperm. Various surface changes have been noted during in
vitro capacitation of mammalian sperm (13, 27). In particular,
the tail of guinea pig sperm demonstrates changes in intra-
membrane particle distribution (28–30) and in lectin agglutin-
ability (31). Migration of the PT-1 antigen into a region where
it was not previously detected means that the antigen is
temporally entering a new environment. Contact between the
PT-1 antigen and anterior tail molecules, not present in the
posterior tail, could result in the activation of new functions.
A precedent for such a model exists in the well-studied acti-
vation of the catalytic unit of adenylate cyclase by contact
with a receptor protein resulting from lateral mobility of
membrane molecules (32). This type of enzyme activation (or
a similar type of inhibition) could occur when the PT-1
antigen gains access to the anterior tail region. Alternatively,
the migration may simply result in the concentration of a
membrane protein in a new surface region where its activity
could have a different effect on cellular function.

The migration of the PH-20 antigen alters the composition
of both the IAM and the posterior head plasma membrane and
occurs immediately after the acrosome reaction. It has
been reported that acrosome reaction is required for guinea
pig sperm to bind to the zona pellucida (14) and acrosome
reaction is known in numerous mammalian species to be
required for sperm-egg plasma membrane fusion (14). PH-20
antigen migration might function in one or both of these steps
in fertilization. On the IAM, the migrated PH-20 antigen
might have a role in zona binding. On the posterior head
surface, a fusion activity could be activated after migration, if
the PH-20 antigen acts in its initial region as an inhibitor.

Control of specific membrane activities through migration
of pre-existing cell surface molecules into or out of cell surface
domains may be important in the regulation of cell surface
function. These rearrangements have the potential to regu-
late cell surface function in response to external stimuli as well
as in response to an internal program during differentiation.

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TABLE I
Inhibition of Migration of the PH-20 Antigen under Various
Conditions

| Condition                  | % Acrosome reaction | % PH-20 Migration |
|----------------------------|---------------------|-------------------|
| No addition                | 100                 | 100               |
| 1 mM CCCP                  | 90                  | 94                |
| 10 mM 2-deoxyglucose       | 100                 | 100               |
| 1 mM CCCP + 10 mM 2-deoxy-
  glucose                   | 100                 | 99                |
| Goat FITC-F(ab')2 anti-mouse IgG | 100           | 0                 |

Sperm, labeled with PH-20 MAb, were resuspended in Ca2+ HEPES with the
inhibitors shown. In the last condition, sperm labeled with PH-20 MAb and
goat FITC-F(ab')2 anti-mouse IgG were resuspended in Ca2+ HEPES. The
samples were incubated at 37°C, 1 μg/ml A23187 was added, and the cells
were fixed 15 min later or 2 h later in the goat F(ab')2 anti-mouse experiment.
Fixed cells were scored for acrosome reaction and fluorescence present on
the IAM and not on the posterior head.

The redistribution of both PT-1 and PH-20 antigens on the
sperm surface resulted from a lateral migration of pre-existing
surface molecules in the absence of added ligands. However,
different mechanisms may have been responsible for the
migration of the PT-1 antigen in contrast to the PH-20
antigen. Before migration, the PT-1 antigen was freely diffus-
ing in the posterior tail region, even though it did not leave
this surface domain. The diffusion coefficient of the PT-1
antigen (2.5 × 10−9 cm2/s), measured by fluorescence redis-
tribution after photobleaching (20), is near the theoretical
limit of a freely diffusing protein in a lipid bilayer. The
mechanism that restricts the PT-1 antigen initially to the
posterior tail region is not known. In this study, we have
shown that PT-1 antigen migration is temperature-dependent,
occurs in the absence of exogenous Ca2+, requires energy, and
is prevented when sperm swimming is prevented. Several
models that would explain these results can be envisioned.
Since the PT-1 antigen is already freely diffusing on the
posterior tail, there may be a barrier in the membrane that
initially prevents the migration of the antigen onto the ante-
or tail region. The antigen may be translocated across this
barrier by an energy-requiring process. Alternatively, energy
may be used to disrupt the barrier or modify the antigen so
as to allow the PT-1 antigen to passively diffuse into the
anterior tail region. In this case, accumulation of the PT-1
antigen in the new region could be explained by trapping of
the antigen.

No migration of the PT-1 antigen occurred when sperm
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