Capacitation of mouse spermatozoa

II. Protein tyrosine phosphorylation and capacitation are regulated by a cAMP-dependent pathway

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SUMMARY

In the accompanying report (Visconti, P.E., Bailey, J.L., Moore, G.D., Pan, D., Olds-Clarke, P. and Kopf, G.S. (1995) Development, 121, 1129-1137) we demonstrated that the tyrosine phosphorylation of a subset of mouse sperm proteins of Mr 40,000-120,000 was correlated with the capacitation state of the sperm. The mechanism by which protein tyrosine phosphorylation is regulated in sperm during this process is the subject of this report. Cauda epididymal sperm, when incubated in media devoid of NaHCO₃, CaCl₂ or bovine serum albumin do not display the capacitation-associated increases in protein tyrosine phosphorylation of this subset of proteins. This NaHCO₃, CaCl₂ or bovine serum albumin requirement for protein tyrosine phosphorylation can be completely overcome by the addition of biologically active, but not inactive, cAMP analogues. Addition of the active cAMP analogues to sperm incubated in media devoid of NaHCO₃, CaCl₂ or bovine serum albumin overcomes the inability of these media to support capacitation, as assessed by the ability of the cells to undergo the zona pellucida-induced acrosome reaction and, in some cases, to fertilize metaphase II-arrested eggs in vitro. The effects of the cAMP analogues to enhance protein tyrosine phosphorylation and to promote capacitation appear to be at the level of the cAMP-dependent protein kinase (PKA), since two specific inhibitors of this enzyme (H-89 and Rp-cAMPS) block the capacitation-dependent increases in protein tyrosine phosphorylation in sperm incubated in media supporting capacitation. Capacitation, as assessed by the aforementioned endpoints, also appears to be inhibited by H-89 in a concentration-dependent manner. These results provide further evidence for the interrelationship between protein tyrosine phosphorylation and the appearance of the capacitated state in mouse sperm. They also demonstrate that both protein tyrosine phosphorylation and capacitation appear to be regulated by cAMP/PKA. Up-regulation of protein tyrosine phosphorylation by cAMP/PKA in sperm is, to our knowledge, the first demonstration of such an interrelationship between tyrosine kinase/phosphatase and PKA signaling pathways.

Key words: mouse, sperm, capacitation, fertilization, in vitro fertilization, tyrosine phosphorylation, cyclic AMP, protein kinase A

INTRODUCTION

Sperm capacitation is a poorly understood maturational process, which occurs in vivo in the female reproductive tract or can be accomplished in vitro in defined media, the endpoint of which confers upon the sperm the ability to undergo the acrosome reaction and to fertilize an egg (Yanagimachi, 1994). Capacitation has been shown to be correlated with changes in sperm plasma membrane fluidity, intracellular ion concentrations, metabolism, and motility (Yanagimachi, 1994 and references therein), but it is not clear how such changes are necessarily tied to the development of this fertilization-competent state. Moreover, although these changes have been known for many years, the molecular basis underlying these events is poorly understood. It is clear from studies in somatic cells that many of these aforementioned cellular changes can be modulated by extrinsic factors that function intracellularly via receptors and second messengers. However, it appears that the initiation and completion of capacitation in mammalian sperm may be regulated by processes inherent to the sperm itself. The mechanism by which such an inherent regulation occurs is not clear, although it is likely that the controlling factor(s) lies with the sperm plasma membrane itself and that changes in the properties of the membrane might lead to a de-repression of
the preprogrammed cellular events that are ultimately necessary for the development of the capacitated state.

We have shown in the preceding report that the acquisition of the capacitated state in caudal epididymal mouse sperm, as assessed by the ability of the cells to acquire the pattern B chlortetracycline fluorescence, to undergo the ZP-induced acrosome reaction, and to fertilize metaphase II-arrested eggs in vitro, is accompanied by the protein tyrosine phosphorylation of a specific subset of sperm proteins (Visconti et al., 1995). The onset of both capacitation and the protein tyrosine phosphorylations are dependent on the presence of serum albumin, Ca\(^{2+}\) and NaHCO\(_3\) in the culture medium, three components that have been demonstrated to be essential for in vitro capacitation of sperm from a variety of different mammals (Yanagimachi, 1994 and references therein). Caput epididymal sperm, which do not possess the ability to undergo capacitation, do not display the protein tyrosine phosphorylations of this subset of proteins. We believe that protein tyrosine phosphorylation may represent an important pathway that may ultimately regulate events associated with capacitation.

The mechanism by which these protein tyrosine phosphorylations are regulated is the subject of this study. Of potential relevance to the requirement of extracellular Ca\(^{2+}\) and NaHCO\(_3\) for these protein tyrosine phosphorylations is the observation that these particular ions have been demonstrated to be involved in the regulation of mammalian sperm cAMP metabolism, and that changes in the concentration of this intracellular second messenger have been linked to both capacitation and the acrosome reaction. For example, it is known that Ca\(^{2+}\)-dependent elevations of guinea pig sperm cAMP concentrations are dependent on the presence of extracellular NaHCO\(_3\) and that the effects of NaHCO\(_3\) on this process do not appear related to changes in intracellular pH (Garbers et al., 1982). The effects of cation and anion may be at the level of the adenylyl cyclase, since the mammalian sperm enzyme has been demonstrated to be regulated directly/indirectly by Ca\(^{2+}\) (Hyne and Garbers, 1979), calmodulin (Gross et al., 1987), and NaHCO\(_3\) (Garty and Salomon, 1987; Okamura et al., 1988; Visconti et al., 1990). Ca\(^{2+}\) and NaHCO\(_3\) also appear to be required for capacitation (Fraser, 1987; Neill and Olds-Clarke, 1987; Boatman and Robbins, 1991) and the induction of the acrosome reaction (Hyne, 1984; Lee and Storey, 1986; Kopf and Gerton, 1991 and references therein; Spira and Breitbart, 1992). Moreover, changes in the activity of adenylyl cyclase and cAMP have been postulated in some species to play a role in capacitation (Perreault and Rogers, 1982; Berger and Clegg, 1983; Boatman and Bavister, 1984; Stein and Fraser, 1984; Monks et al., 1986; Yanagimachi, 1994 and references therein).

The interrelationships between NaHCO\(_3\), Ca\(^{2+}\), cAMP, protein tyrosine phosphorylation and capacitation in cauda epididymal mouse sperm was further examined in the present report. The capacitation-associated changes in protein tyrosine phosphorylation that are dependent on the presence of bovine serum albumin (BSA), Ca\(^{2+}\) and HCO\(_3^-\) can be mimicked in the absence of each of these required media constituents by the addition of active, but not inactive, membrane-permeable cAMP analogues. Correlated with the stimulatory effects of these cAMP analogues on protein tyrosine phosphorylation is a stimulatory effect of these analogues on capacitation. The stimulatory effects of cAMP on protein tyrosine phosphorylation and capacitation appear to be at the level of PKA.

**MATERIALS AND METHODS**

**Reagents**

Sp-cAMPS, Rp-cAMPS, 8-bromo cAMP and KN-62 were from Research Biochemicals International (Natick, MA). H89 (N-[2-(p-bromocinnamylamino)ethyl]-5-isouquinolinesulfonamide), staurosporine and chelerythrine were from LC Labs (Woburn, MA). Dibutyryl cAMP and IBMX were purchased from Sigma. Anti-phosphotyrosine antibody (clone 4G10) was obtained from UBI (Placid, NY). All other reagents were of the highest analytical grade.

**Culture media**

The basic medium used throughout these studies for the preparation of sperm was a modified Krebs Ringer bicarbonate medium (HMB-Hepes buffered), as described by Lee and Storey (1986). The preparation of the media is as described in the accompanying report (Visconti et al., 1995).

**Preparation of sperm and in vitro fertilization**

Uncapacitated caudal epididymal sperm were collected from CD1 retired breeder male mice by the methods described in the accompanying paper (Visconti et al., 1995). In vitro fertilization of metaphase II-arrested eggs was performed as previously described (Moore et al., 1993), with the modifications described in the accompanying report (Visconti et al., 1995).

**Preparation and solubilization of mouse zonae pellucidae**

Zonae pellucidae were prepared from homogenized oocytes of virgin female 22-day old outbred CD-1 mice (Charles River) as described in Ward et al. (1992) and solubilized for all experiments by the procedures outlined in the previous report (Visconti et al., 1995).

**Chlortetracycline assay**

This assay was used to assess both the capacitation state of the sperm and the percentage of acrosome reactions. The experimental design and use of these assays is described in Visconti et al. (1995).

**Preparation of sperm membranes and measurement of adenylyl cyclase activity**

Caudal epididymal spermatozoa were recovered as described by Endo et al. (1987, 1988), and incubated at a concentration of 1-3x10^8 cells/ml for 1 hour at 37°C in a humidified incubator with 5% CO\(_2\) in air in Hepes-buffered modified bicarbonate Krebs-Ringer medium (HMB) containing 20 mg/ml BSA. At the end of the incubation, the sperm cells were diluted twice in HMB+BSA and centrifuged for 20 minutes (150 g) at room temperature. The resultant pellet was resuspended in ice-cold homogenization buffer (50 mM Hepes, 1 mM EGTA, 10% DMSO, pH 7.5) and gently homogenized (10 strokes in a loose glass-teflon homogenizer) under conditions that have been shown to maintain the ZP-G protein coupling (Ward et al., 1992). The homogenate was subjected to a low speed centrifugation (1,000 g; 4°C) to pellet the membranes. The particulate crude membrane fraction was resuspended in the same homogenization buffer, aliquoted and kept frozen (~70°C). The adenylyl cyclase assays were performed on membrane preparations less than a week old. Generally, 250-350 μg of membrane proteins were obtained using 20 males. Adenylyl cyclase activity was measured by the conversion of ([α-^32P]ATP to [α-^32P]cAMP (Salomon et al., 1974). The assay was carried out for 20 minutes at 37°C in the presence of 50 mM Hepes, 1.5 mM ATP, 15 mM MgCl\(_2\), 10 mM KCl, 4 mM DTT, 1 mM IBMX, 16 μg creatine kinase, 3.2 mM creatine phosphate and 2.5-3 μCi ([α-
Cyclic AMP analogues can substitute for bicarbonate anion in supporting both protein tyrosine phosphorylation and capacitation

We first examined whether the mouse sperm adenyl cyclase could be stimulated by HCO$_3^-$ ions. Membranes from capacitated mouse sperm were prepared under conditions that support ZP3-AR spont. Membranes from capacitated mouse sperm were prepared and assayed for adenyl cyclase activity using Mg$^{2+}$ as the divalent cation as described in Materials and Methods. The final concentration of NaHCO$_3$ was adjusted to different concentrations and assayed for adenylyl cyclase activity in the absence and presence of various concentrations of NaHCO$_3$. As shown in Fig. 1, NaHCO$_3$ stimulated adenyl cyclase activity in a concentration-dependent manner that was maximal at 50 mM. These stimulatory effects compare favorably with those effects seen by Okamura et al. (1985), Garty and Salomon (1987) and Visconti et al. (1990). The effect of NaHCO$_3$ on mouse sperm adenyl cyclase activity is due to the effect of HCO$_3^-$ ions, since substituting the NaHCO$_3$ with NaCl had no effect on the activity of the enzyme (data not shown). Moreover, the effect of NaHCO$_3$ on enzyme activity did not result as a consequence of changes in assay pH, since adjusting the pH of the buffers used in the enzyme assay from pH 7-8 had no effect on enzyme activity (data not shown).

Cyclic AMP analogues can substitute for bicarbonate anion in supporting both protein tyrosine phosphorylation and capacitation

As previously demonstrated, the presence of NaHCO$_3$ in the medium is required for capacitation and the associated changes in protein tyrosine phosphorylation (Visconti et al., 1995, Figs 6, 7). Since NaHCO$_3$ stimulated the mouse sperm adenyl cyclase (Fig. 1), we determined whether the addition of membrane permeable cAMP analogues could substitute for NaHCO$_3$ in promoting both protein tyrosine phosphorylation and capacitation. Caudal epididymal sperm were incubated for different periods of time in complete medium or in medium devoid of NaHCO$_3$ in either the absence or presence of 1 mM dibutyryl cAMP (db-cA) plus 100 µM IBMX. When compared to sperm incubated in complete medium, sperm incubated in medium devoid of NaHCO$_3$ did not display the capacitation-associated changes in protein tyrosine phosphorylation, even after incubation times up to 5 hours (Fig. 2A). However, when the NaHCO$_3$-free culture medium was supplemented with 1 mM db-cA plus 100 µM IBMX, the protein tyrosine phosphorylation was observed and this was maximal by 3 hours of incubation (Fig. 2A). As described in the previous report (Visconti et al., 1995), the degree of tyrosine phosphorylation of the phosphotyrosine-containing protein in mouse sperm membranes (Kalab et al., 1994), was not affected by the absence of NaHCO$_3$ in the medium and its phosphorylation did not appear to be modulated by the addition of 1 mM db-cA plus 100 µM IBMX (Fig. 2A). These effects on protein tyrosine phosphorylation were not limited to db-cA since 1 mM 8 bromo-cAMP (8Br-cA) and 1 mM Sp-cAMPS (Sp-cA) also supported protein tyrosine phosphorylation in the absence of NaHCO$_3$ and in the presence of IBMX (Fig. 2B, right-hand panel). In contrast to the effects of these active cAMP analogues, addition of Rp-cAMPS (Rp-cA), which is a membrane-permeable cAMP analogue that binds to the regulatory subunit of PKA but fails to cause subunit dissociation and activation of PKA and thus antagonizes CAMP action (Van Haastert et al., 1984; Parker Botelho et al., 1988), did not support protein tyrosine phosphorylation in the absence of NaHCO$_3$ (Fig. 2B, right-hand panel). In contrast to the effects seen when 8Br-cA was added to the sperm in the presence of IBMX, 8Br-cA added in the absence of IBMX failed to efficiently support protein tyrosine phosphorylation (Fig. 2B, left-
hand panel). Such a differential effect was not observed when Sp-cA was used (Fig. 2B, compare left-hand and right-hand panels); this is probably due to its inability to be hydrolyzed by cyclic nucleotide phosphodiesterases. These effects appeared to be specific for cAMP analogues since dibutyryl cGMP, in either the absence or presence of IBMX, had no effect on protein tyrosine phosphorylation (data not shown).

Sperm incubated in medium devoid of NaHCO₃ also did not display the CTC fluorescence B pattern indicative of the capacitated state (Fig. 3A). The ZP-induced acrosome reaction was, P. E. Visconti and others

Fig. 2. Effects of cyclic AMP analogues and 3-isobutyl-1-methylxanthine (IBMX) on the appearance of phosphotyrosine-containing proteins in caudal epididymal mouse sperm incubated in media devoid of NaHCO₃. (A) Sperm were incubated for the periods of time indicated in the figure in HM medium (no added NaHCO₃) or in the same medium containing 1 mM dibutyryl cAMP (db-cA) plus 100 µM IBMX. At those times, an aliquot of the sperm suspension was removed (5×10⁵ sperm), centrifuged and the whole sperm extract subjected to SDS-PAGE by the procedures outlined in Materials and Methods. The proteins were then transferred to Immobilon P and probed with a monoclonal antibody to phosphotyrosine (α pY). This experiment was performed at least three times with similar results. Shown is a representative experiment. (B) Sperm were incubated in NaHCO₃-depleted medium (−), or in NaHCO₃-depleted medium containing 1 mM Sp-cAMPS (Sp-cA), Rp-cAMPS (Rp-cA) or 8 bromo-cAMP (8Br-cA) in either the absence (left-hand panel) or presence (right-hand panel) of 100 µM IBMX. This experiment was performed at least three times with similar results; a representative experiment is shown. In both A and B, the major phosphotyrosine-containing protein that appears at Mr 116,000 in all of the lanes is the p95/116 hexokinase. Numbers on the left-hand side of this gel and in subsequent figures represent the position of the relative molecular mass standards (×10⁻³).

Fig. 3. Effects of dibutyryl cAMP plus 3-isobutyl-1-methylxanthine (IBMX) on the percentage of caudal epididymal mouse sperm displaying chlortetracycline fluorescence pattern ‘B’ and the percentage of sperm undergoing the spontaneous and zona pellucida-induced acrosome reaction (chlortetracycline pattern ‘AR’) in media containing NaHCO₃ or in media devoid of NaHCO₃. Sperm were incubated for a period of 90 minutes in NaHCO₃-free HMB, NaHCO₃-free HMB containing 10 mM NaHCO₃, or in these media containing 1 mM dibutyryl cAMP (db-cA) plus 100 µM IBMX. (A) Sperm were assessed for the percentage of CTC pattern ‘B’, indicative of the capacitated state. Data represent the mean ± s.e.m., n=3. (B) Sperm were assessed for the percentage of CTC pattern ‘AR’, representing cells undergoing acrosomal exocytosis, following a 30 minute incubation in buffer [open bars] or solubilized zona pellucidae (5 ZP/µl) [solid bars]. Data represent the mean ± s.e.m., n=3. Statistical analysis was performed as described in Materials and Methods; *P<0.05 and **P<0.025.
likewise, completely dependent on the presence of NaHCO$_3$ in the medium (Fig. 3B). As was seen with protein tyrosine phosphorylation, the addition of 1 mM db-cA plus 100 µM IBMX to medium devoid of NaHCO$_3$ could substitute for NaHCO$_3$ in supporting the ability of the sperm to display the CTC fluorescence B pattern and to undergo the ZP-induced acrosome reaction (Fig. 3A,B).

**Cyclic AMP analogues can substitute for Ca$^{2+}$ in supporting both protein tyrosine phosphorylation and capacitation**

When compared to complete medium, sperm incubated in medium devoid of added Ca$^{2+}$ failed to display the capacitation-dependent changes in protein tyrosine phosphorylation (Fig. 4; Visconti et al., 1995). When sperm were incubated in Ca$^{2+}$-free medium containing various cAMP analogues plus IBMX for a period of 90 minutes, the capacitation-dependent changes in protein tyrosine phosphorylation were observed when biologically active (Sp-cA; db-cA; 8Br-cA), but not inactive (Rp-cA), cAMP analogues were used (Fig. 4, right-hand panel). It is of interest that IBMX alone supported protein tyrosine phosphorylation, but not to the degree of that observed in the presence of the active cAMP analogues (Fig. 4, right-hand panel). This is due, presumably, to its function as an inhibitor of cyclic nucleotide phosphodiesterase activity. IBMX alone also supported a greater degree of protein tyrosine phosphorylation but Rp-cA appeared to inhibit the IBMX-induced changes in protein tyrosine phosphorylation. This is presumably due to the fact that Rp-cA, in addition to its ability to bind to the regulatory subunit of PKA but not activate the enzyme, has been demonstrated to function as a cAMP antagonist (Van Haastert et al., 1984; Parker Botelho et al., 1988) and might therefore abrogate the effects of IBMX as a cyclic nucleotide phosphodiesterase inhibitor by competing with endogenous cAMP. In contrast to the effects seen in the presence of IBMX, in the absence of this phosphodiesterase inhibitor only Sp-cA appeared to support the appearance of the capacitation-dependent changes in protein tyrosine phosphorylation (Fig. 4, left-hand panel). This is presumably due to the fact that this analogue is very effective in activating PKA and is very poorly hydrolyzed by cyclic nucleotide phosphodiesterases. These data suggest that cAMP analogues can substitute for Ca$^{2+}$ deficiency in supporting protein tyrosine phosphorylation.

Sperm incubated in medium devoid of Ca$^{2+}$, likewise, did not display the CTC fluorescence B pattern indicative of the capacitated state (Fig. 5A). The ZP-induced acrosome reaction was also completely dependent on the presence of Ca$^{2+}$ in the medium (Fig. 5B). As was seen with protein tyrosine phosphorylation, the addition of 1 mM db-cA plus 100 µM IBMX to medium devoid of Ca$^{2+}$ could substitute for Ca$^{2+}$ in supporting the ability of the sperm to display the CTC fluorescence B pattern (Fig. 5A). In contrast, these agents did not support the ability of the sperm to undergo a ZP-induced acrosome reaction (Fig. 5B), presumably due to the fact that extracellular Ca$^{2+}$ is essential for this exocytotic event (Yanagimachi, 1994).

**Cyclic AMP analogues can substitute for serum albumin in supporting both protein tyrosine phosphorylation and capacitation**

Since serum albumin is required for both capacitation (Yanagimachi, 1994 and references therein) and the appearance of the phosphorytrosine-containing proteins in caudal epididymal mouse sperm incubated in media devoid of CaCl$_2$. Sperm were incubated for 90 minutes in HMB medium containing no added CaCl$_2$ (Ca$^{2+}$-free HMB) in either the absence (left-hand panel) or presence (right-hand panel) of 100 µM IBMX. These media were also supplemented with either no cAMP analogues (--) or with 1 mM Rp-cAMPS (Rp-cA), 1 mM Sp-cAMPS (Sp-cA), 1 mM dibutylcAMP (db-cA) or 0.5 mM 8-bromo-cAMP (8Br-cA). Sperm were also incubated in the presence of complete HMB (extreme right lane marked +Ca$^{2+}$). This experiment was performed at least three times with similar results. Shown, is a representative experiment.

![Fig. 4. Effects of cyclic AMP analogues and 3-isobutyl-1-methylxanthine (IBMX) on the appearance of phosphorytrosine-containing proteins in caudal epididymal mouse sperm incubated in media devoid of CaCl$_2$. Sperm were incubated for 90 minutes in HMB medium containing no added CaCl$_2$ (Ca$^{2+}$-free HMB) in either the absence (left-hand panel) or presence (right-hand panel) of 100 µM IBMX. These media were also supplemented with either no cAMP analogues (--) or with 1 mM Rp-cAMPS (Rp-cA), 1 mM Sp-cAMPS (Sp-cA), 1 mM dibutylcAMP (db-cA) or 0.5 mM 8-bromo-cAMP (8Br-cA). Sperm were also incubated in the presence of complete HMB (extreme right lane marked +Ca$^{2+}$). This experiment was performed at least three times with similar results. Shown, is a representative experiment.](image-url)
Gerton, 1991; Tash et al., 1988) that may be more active in sperm incubated in the presence of Ca^{2+}. In contrast to the effects seen in the presence of IBMX, in the absence of this phosphodiesterase inhibitor only Sp-cA appeared to support the spontaneous and zona pellucida-induced acrosome reaction (chlorotetracycline pattern ‘AR’) in medium containing CaCl_2 or in medium devoid of CaCl_2. Sperm were incubated for a period of 90 minutes in Ca^{2+}-free HMB, Ca^{2+}-free HMB containing 1.7 mM CaCl_2, or in these media containing 1 mM dibutyryl cAMP (db-cA) plus 100 µM IBMX.

Sperm incubated in medium devoid of BSA, likewise, did not display the CTC fluorescence B pattern indicative of the capacitated state (Fig. 7A). Capacitation was also not seen in sperm incubated in medium containing 1 mM db-cA plus 100 µM IBMX (Fig. 7B). The addition of 1 mM db-cA plus 100 µM IBMX to the BSA-free medium, however, could substitute fully for BSA in supporting the ability of the sperm to display the CTC fluorescence B pattern (Fig. 7A) but not to undergo the ZP-induced acrosome reaction (Fig. 7B).

Although the effects of BSA depletion on capacitation using in vitro fertilization as an endpoint could not be assessed due to the fact that serum albumin is needed for zygote formation, we observed that sperm incubated under capacitation conditions in the absence of BSA but in the presence of 1 mM db-cA plus 100 µM IBMX, and then incubated with ZP-intact eggs in the presence of BSA, displayed a higher percentage fertilization (79%; 32 eggs assessed) than sperm incubated in capacitation medium devoid of both BSA and db-cA/100 µM IBMX (55%; 48 eggs assessed) (data not shown). These data are consistent with the notion that db-cA plus IBMX may function in the absence of BSA to accelerate the capacitation process and/or synchronize capacitation in a greater population of sperm (Fraser, 1981).
Protein tyrosine phosphorylation is accelerated by dibutyryl cAMP and IBMX in media that support capacitation

Since it has been demonstrated that dibutyryl cAMP accelerates capacitation in vitro in mouse sperm (Fraser, 1981), we examined whether the addition of db-cA plus IBMX influenced the time-dependent onset of protein tyrosine phosphorylation that occurs during the normal time course of capacitation in vitro. As shown in Fig. 8, when compared to controls, addition of 1 mM db-cA plus 100 µM IBMX to sperm in complete medium significantly reduced the time during which changes in protein tyrosine phosphorylation occurred.

Protein tyrosine phosphorylation and capacitation appear to be regulated by cAMP-dependent protein kinase

Since the primary intracellular target for cAMP action is the PKA, we examined whether PKA played a role in protein tyrosine phosphorylation and capacitation. Caudal epididymal sperm were incubated for 90 minutes in complete medium that supports capacitation in either the absence or presence of N-[2-(p-bromocinnamylamino)ethyl]-5-isoquinolinesulfonamide (H-89), a highly selective and cell permeable inhibitor of PKA (Chijiwa et al., 1990). As shown in Fig. 9A, H-89 inhibited protein tyrosine phosphorylation of sperm proteins in a concentration-dependent manner with maximal effects observed at 10 µM. In contrast, the tyrosine phosphorylation of the p95/116 hexokinase was unaffected by H-89 at all concentrations tested. In contrast to the effects of this PKA inhibitor on protein tyrosine phosphorylation, inhibitors of protein kinase C (1 µM staurosporine; 10 µM chelerythrine) and Ca²⁺/calmodulin-dependent protein kinase (10 µM KN-62) had no effects on protein tyrosine phosphorylation that accompanied capacitation (data not shown).

Sperm incubated in complete medium for 90 minutes did not display protein tyrosine phosphorylation when incubated with the cAMP antagonist Rp-cAMPS, whereas phosphorylation was observed in either the absence of a cAMP analogue or in the presence of polyvinyl alcohol. Sperm were incubated for a period of 90 minutes in BSA-free HMB, BSA-free HMB containing 1 mg/ml polyvinyl alcohol (PVA), HMB containing 3 mg/ml BSA, or in these media containing 1 mM dibutyryl cAMP (db-cA) plus 100 µM IBMX. (A) Sperm were assessed for the percentage of CTC pattern ‘B’, indicative of the capacitated state. Data represent the mean ± s.e.m., n=3. (B) Sperm were assessed for the percentage of CTC pattern ‘AR’, representing cells undergoing acrosomal exocytosis, following a 30 minute incubation in buffer (open bars) or solubilized zona pellucidae (5 ZP/µl; closed bars). Data represent the mean ± s.e.m., n=3. Statistical analysis was performed as described in Materials and Methods; *P<0.01, **P<0.025 and ***P<0.005.

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Sperm incubated in complete medium for 90 minutes did not display protein tyrosine phosphorylation when incubated with
H-89 was also observed to inhibit capacitation, as assessed by a number of criteria. The appearance of the CTC B pattern of fluorescence was inhibited by H-89 in a concentration-dependent manner (Fig. 10A). This was accompanied by a loss in the ability of the sperm to undergo a ZP-induced acrosome reaction (Fig. 10B). Moreover, if sperm were incubated under capacitating conditions in the presence of 30 µM H-89, a concentration that inhibits tyrosine phosphorylation, the appearance of the CTC B pattern of fluorescence and the ZP-induced acrosome reaction (Figs 9, 10), in vitro fertilization of ZP-intact eggs is inhibited, as evidenced by a significant reduction in the formation of a pronucleus in eggs inseminated with H-89-treated sperm (Fig. 11, H-89 in capacitation media; H-89 in both capacitation and fertilization media). The effect on fertilization is especially pronounced when the H-89 concentration is held at 30 µM by keeping the inhibitor in both the capacitation and fertilization media. The reduction in the inhibitory effect on fertilization observed when the H-89 is present only in the capacitation medium is probably due to the dilution of the H-89 at the time of in vitro insemination (from 30 µM to 0.2 µM). The lowering of the H-89 concentration during the 2 h insemination period might then permit capacitation to occur so that the sperm can bind to the ZP, undergo the acrosome reaction, penetrate and fertilize the egg. Consistent with this hypothesis is the fact that after dilution of the sperm to the same extent in an H89-free medium, the protein tyrosine phosphorylation increases steadily (data not shown). The inhibitory effects of H-89 on fertilization does not appear to be due to an inhibitory effect of H-89 on the egg itself, since insemination of ZP-free eggs in H89-containing fertilization medium with sperm that have been capacitated in the absence of H-89 does not inhibit in vitro fertilization (data not shown).

**DISCUSSION**

The data outlined in this report further support the interrelationship between protein tyrosine phosphorylation and the capacitation state of mouse sperm, since modulation of capacitation by the addition of cAMP analogues also modulates protein tyrosine phosphorylation. Moreover, inhibition of protein tyrosine phosphorylation and capacitation by two inhibitors of PKA (Rp-cAMP; H89) that act on this enzyme by totally different mechanisms further support the linkage between these two events and strengthen the argument that PKA is required for the regulation of these two processes. The

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**Fig. 9.** Effects of the protein kinase A inhibitor H-89, as well as Rp-cAMPS and Sp-cAMPS, on the appearance of phosphotyrosine-containing proteins in caudal epididymal mouse sperm incubated in complete media that support capacitation. (A) Sperm were incubated for 90 minutes in complete HMB medium in the absence or presence of increasing concentrations of H-89 and protein tyrosine phosphorylation assessed. (B) Sperm were incubated for 90 minutes in complete HMB medium in either the absence (-) or presence of 1 mM Rp-cAMPS (Rp-cA), 1 mM Sp-cAMPS (Sp-cA) or 30 µM H-89, and protein tyrosine phosphorylation assessed. This experiment was performed at least three times with similar results. Shown, is a representative experiment.

H-89 was also observed to inhibit capacitation, as assessed by a number of criteria. The appearance of the CTC B pattern fluorescence was inhibited by H-89 in a concentration-dependent manner (Fig. 10A). This was accompanied by a loss in the ability of the sperm to undergo a ZP-induced acrosome reaction (Fig. 10B). Moreover, if sperm were incubated under capacitating conditions in the presence of 30 µM H-89, a concentration that inhibits tyrosine phosphorylation, the appearance of the CTC B pattern of fluorescence and the ZP-induced acrosome reaction (Figs 9, 10), in vitro fertilization of ZP-intact eggs is inhibited, as evidenced by a significant reduction in the formation of a pronucleus in eggs inseminated with H-89-treated sperm (Fig. 11, H-89 in capacitation media; H-89 in both capacitation and fertilization media). The effect on fertilization is especially pronounced when the H-89 concentration is held at 30 µM by keeping the inhibitor in both the capacitation and fertilization media. The reduction in the inhibitory effect on fertilization observed when the H-89 is present only in the capacitation medium is probably due to the dilution of the H-89 at the time of in vitro insemination (from 30 µM to 0.2 µM). The lowering of the H-89 concentration during the 2 h insemination period might then permit capacitation to occur so that the sperm can bind to the ZP, undergo the acrosome reaction, penetrate and fertilize the egg. Consistent with this hypothesis is the fact that after dilution of the sperm to the same extent in an H89-free medium, the protein tyrosine phosphorylation increases steadily (data not shown). The inhibitory effects of H-89 on fertilization does not appear to be due to an inhibitory effect of H-89 on the egg itself, since insemination of ZP-free eggs in H89-containing fertilization medium with sperm that have been capacitated in the absence of H-89 does not inhibit in vitro fertilization (data not shown).

**DISCUSSION**

The data outlined in this report further support the interrelationship between protein tyrosine phosphorylation and the capacitation state of mouse sperm, since modulation of capacitation by the addition of cAMP analogues also modulates protein tyrosine phosphorylation. Moreover, inhibition of protein tyrosine phosphorylation and capacitation by two inhibitors of PKA (Rp-cAMPS; H89) that act on this enzyme by totally different mechanisms further support the linkage between these two events and strengthen the argument that PKA is required for the regulation of these two processes. The
The fact that biologically active cAMP analogues can overcome the deficiency in protein tyrosine phosphorylation and capacitation seen in media devoid of NaHCO3, Ca2+ or BSA suggests that this second messenger functions downstream from events stimulated by these media constituents. Although some cAMP analogues added alone could completely substitute for specific media components in supporting protein tyrosine phosphorylation, the effects of some of these analogues on protein tyrosine phosphorylation was augmented when the experiments were carried out in the presence of H-89 and then diluted into fertilization medium devoid of H-89, the final H-89 concentration in the insemination droplet was 0.2 µM. Following insemination and washing the eggs were then evaluated for the presence of a pronucleus. Numbers in parentheses above the bars indicate the numbers of eggs analyzed in the different treatment groups. The data represent the mean ± s.e.m. of three individual experiments. Statistical analysis was performed as described in Materials and Methods; *P<0.01, **P<0.005.

Fig. 11. Effects of the protein kinase A inhibitor H-89 on in vitro fertilization of zona pellucida-intact and zona pellucida-free metaphase II-arrested eggs. Sperm were capacitated in either the absence (−) or presence (+) of 30 μM H-89 for a period of 2 hours and then incubated with zona pellucida-intact eggs for 2 hours in fertilization medium containing no added H-89 (−) or 30 μM H-89 (+), according to the procedures outlined in the Methods. When sperm were capacitated in the presence of H-89 and then diluted into fertilization medium devoid of H-89, the final H-89 concentration in the insemination droplet was 0.2 µM. Following insemination and washing the eggs were then evaluated for the presence of a pronucleus. Numbers in parentheses above the bars indicate the numbers of eggs analyzed in the different treatment groups. The data represent the mean ± s.e.m. of three individual experiments. Statistical analysis was performed as described in Materials and Methods; *P<0.01, **P<0.005.

As discussed in the accompanying manuscript (Visconti et al., 1995), it is of interest that the sperm appears to intrinsically control events leading to capacitation and that the controlling factor(s) related to this maturational process may lie within the plasma membrane. Destabilization of the plasma membrane by changes in membrane fluidity could ultimately set in motion a series of intrinsic pre-programmed membrane and intracellular events that ultimately regulate those sperm functions that are normally associated with the capacitated state (e.g., changes in sperm motility; ability to undergo a ZP-induced acrosome reaction). Our data would not only support the idea that protein tyrosine phosphorylation might represent a key set of signaling events that are entrained as part of the onset of the capacitated state but that there appears to be a regulatory hierarchy that involves PKA.

Fig. 12 provides a working model for how such changes at the level of the sperm plasma membrane might engage a Ca2+-dependent pathway that leads to protein tyrosine phosphorylation and the concomitant endpoints of capacitation. In this model, serum albumin, by virtue of its ability to serve as an extracellular sink for sperm membrane cholesterol efflux (Davis, 1976, 1980; Davis et al., 1979a,b; Go and Wolf, 1985; Langlais and Roberts, 1985), could lead to alterations in membrane fluidity with resultant changes in the permeability of the sperm to Ca2+ (Singh et al., 1978; Ruknudin and Silver, 1990) and/or HCO3− (Okamura et al., 1988). Changes in the permeability of the sperm towards these ions can then regulate intracellular cAMP concentrations through stimulatory effects.
on adenylyl cyclase. Adenylyl cyclase could be regulated either directly by Ca\(^{2+}\) (Hyne and Garbers, 1979) or via calmodulin (Kopf and Vacquier, 1984; Gross et al., 1987). Moreover, sperm cAMP concentrations could be additionally modulated by Ca\(^{2+}\) at the level of a Ca\(^{2+}\)/calmodulin-stimulated cyclic nucleotide phosphodiesterase. Adenylyl cyclase could, likewise, be regulated by HCO\(_3\)\(^{-}\), as we have demonstrated in Fig. 1 and has been also demonstrated in sperm from other species (Garty and Salomon, 1987; Okamura et al., 1985; Visconti et al., 1990). The resultant changes in cAMP would modulate the activities of PKA, of which there appears to be both Types I and II present in mouse sperm (Visconti and Kopf, unpublished observations), and such changes in enzyme activity would lead ultimately to changes in protein tyrosine phosphorylation.

The endpoints of capacitation that such tyrosine phosphorylations might regulate are also unknown. Changes in sperm motility appear to accompany capacitation process (Neill and Olds Clarke, 1987; Olds Clarke, 1990). Since it is known that cAMP plays an important regulatory role in the initiation of flagellar motility (Tash and Means, 1983) and in alterations of the flagellar curvature (Lindemann and Kanous, 1989; White and Aitken, 1989; Lindemann et al., 1991; Si and Okuno, 1993), it is possible that such cAMP-dependent changes in protein tyrosine phosphorylation might be related to these motility changes. For example, it has been demonstrated that the initiation of flagellar motility in the sperm of the rainbow trout is accompanied by the tyrosine phosphorylation of a 15,000 M\(_r\) flagellar protein that occurs in a cAMP-dependent manner (Hayashi et al., 1987). This protein and the tyrosine kinase that regulates its phosphorylation have yet to be identified. It is also possible that changes in tyrosine kinase and/or phosphoprotein phosphatase activities and resultant protein tyrosine phosphorylations are involved in the regulation of ion channels (Siegelbaum, 1994 and references therein). It is interesting to note that the regulation of the voltage-gated cation channel in Aplysia bag neurons by tyrosine phosphorylation appears to be modulated in a cAMP-dependent manner (Wilson and Kaczmarek, 1993).

Perhaps the most intriguing observations that have resulted from these studies are that the protein tyrosine phosphorylation state in mouse sperm appears to be regulated by cAMP through the action of PKA. To the best of our knowledge, these observations provide the first example of an up-regulation of protein tyrosine phosphorylation by cAMP/PKA, and suggests that there is cross-talk at some level between PKA and tyrosine kinase/phosphoprotein phosphatase signaling pathways in the sperm. Evidence exists for the down regulation of receptor tyrosine kinase actions by cAMP (Wu et al., 1993; Cook and McCormick, 1993) but up-regulation has thus far not been demonstrated. Protein tyrosine phosphatase activity has been demonstrated to be both inhibited (Begum et al., 1992) and stimulated (Brautigan and Pinault, 1991) by cAMP, effects of which could ultimately modulate the levels of protein tyrosine phosphorylation. The level at which cAMP/PKA functions to regulate the steady state levels of protein tyrosine phosphorylation in mouse sperm under conditions conducive to capacitation (i.e., tyrosine kinases; phosphoprotein phosphatases) will be the subject of intense scrutiny in our laboratory for the foreseeable future. Subsequent investigations into the mechanism by which cAMP/protein kinase regulates those tyrosine kinases/phosphoprotein phosphatases might yield new information about alternative modes of cross-talk between these two signaling pathways.

This work was supported by grants from the NIH to G. S. K. (HD 06274; HD22732) and P. O. C. (HD15045). P. E. V. and G. D. M. were supported by grants from the Rockefeller Foundation. J. L. B. was supported by an NIH grant to Dr Bayard T. Storey (HD06274). P. L. was supported by a fellowship from the Natural Sciences and


(Accepted 3 January 1995)