



Published in final edited form as:

Reproduction. 2024 August 01; 168(2): . doi:10.1530/REP-24-0069.

Dual role of Valosin-Containing Protein (VCP/p97) in mouse sperm during capacitation

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Abstract

Valosin-containing protein (VCP; aka p97), a member of the AAA family (ATPases Associated with various cellular Activities), has been associated with a wide range of cellular functions. While previous evidence has shown its presence in mammalian sperm, our study unveils its function in mouse sperm. Notably, we found that mouse VCP does not undergo tyrosine phosphorylation during capacitation and exhibits distinct localization patterns. In the sperm head, it resides within the equatorial segment and, following exocytosis, it is released and cleaved. In the flagellum, VCP is observed in the principal and midpiece. Furthermore, our research highlights a unique role for VCP in the cAMP/PKA pathway during capacitation. Pharmacological inhibition of sperm VCP led to reduced intracellular cAMP levels that resulted in decreased phosphorylation in PKA substrates and tyrosine residues, and diminished fertilization competence. Our results show that in mouse sperm, VCP plays a pivotal role in regulating cAMP production, probably by the modulation of soluble adenylyl cyclase (sAC) activity.

Keywords

cAMP; PKA; tyrosine phosphorylation; NMS-873; acrosomal exocytosis

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Author contribution statement

MJ, FALS, JSE performed experiments and analysed data, Methodology: MGE and CLMB; Conceptualization: Dario K, Diego K and PEV. GML and MGB conceived the study and wrote the paper.

Introduction

Following ejaculation, sperm are not immediately able to fertilize an egg. To achieve this task, they require to undergo several maturational steps in the female reproductive tract, all grouped under the name of sperm capacitation (Austin, 1951; Chang, 1951). Sperm capacitation involves complex signaling pathways that prepare sperm to develop hyperactivated motility and to undergo acrosomal exocytosis (aka acrosome reaction, AR) (Gervasi and Visconti, 2016; Stival et al., 2016). Because sperm cells are transcriptionally and translationally silent, their regulatory signaling pathways largely depend on changes in intracellular ion concentrations, synthesis and degradation of second messengers, and protein post-translational modifications. The proteins that display either a change in localization or post-translational modification during capacitation are of particular interest because of their potential role in fertilization. Although the capacitation-induced molecular events have been intensively scrutinized in recent decades, many proteins modified during this process remain poorly understood.

Valosin-containing protein (VCP, aka p97) has been reported to undergo tyrosine phosphorylation and relocalization during capacitation in human sperm (Ficarro et al., 2003). VCP is a member of the AAA family (ATPases Associated with various cellular Activities) (Zhang et al., 2000) that has been implicated in a wide variety of cellular functions. It was initially recognized as one of the endoplasmic reticulum (ER) associated proteins, and has been shown to play critical roles in regulating ER formation and morphology by participating in the ubiquitin–proteasome system (UPS) and other intracellular signaling pathways (Shih and Hsueh, 2018). VCP has also been implicated in several fusion events in both yeast and mammalian cells (Kondo et al., 1997). Together with N-ethylmaleimide-sensitive factor (NSF) and syntaxin 5, VCP regulates the fusion of Golgi membranes (Rabouille et al., 1995). Even though VCP and NSF are highly homologous, they appear to participate in different fusion events, likely due to the participation of specific cofactors (Urbé et al., 1998). It was also reported that VCP has a role as a chaperone and aids in the assembly, disassembly, and functional operation of protein complexes (White and Lauring, 2007). During T cell activation, VCP undergoes tyrosine phosphorylation, which does not alter its ATPase activity (Egerton and Samelson, 1994), but regulates its subcellular localization (Madeo et al., 1998). Moreover, tyrosine phosphorylation of VCP is required for membrane fusion processes such as transitional ER assembly *in vitro* (Hirling et al., 1996).

In human sperm, during the capacitation process, along with its phosphorylation on tyrosine residues, VCP changed its localization pattern. In non-capacitated human sperm, VCP localizes to the neck and tail regions. However, upon capacitation, fluorescent staining shows the appearance of VCP in the anterior head. VCP has also been proposed as a substrate of cAMP-activated boar sperm tyrosine kinase (Geussova et al., 2002) and it has been shown to participate in the autophagy and UPS of mitophagy after mammalian fertilization (Song et al., 2016).

In this article, we study the localization and function of VCP in mouse sperm. We show that VCP has a dual role in this species. First, VCP localizes in the equatorial segment (ES) of the sperm head and is cleaved and released after the acrosome reaction. Second,

contrary to what was observed for human, in mouse sperm, VCP does not undergo tyrosine phosphorylation on tyrosine residues during capacitation. Additionally, we found that VCP localizes to the flagellum and is involved in signaling events regulating the cAMP/PKA pathway, whose activation is required for capacitation.

Materials and methods

Reagents

Bovine serum albumin (BSA, A7906), α - β -tubulin (T4026), protease inhibitor cocktail (P8340), and progesterone (P8783) were purchased from Sigma-Aldrich (St. Louis, MO, USA) unless stated otherwise. Chemicals were obtained from the following sources: α -phosphotyrosine (α -pY) clone 4G10 from Millipore (Temecula, CA, USA); horseradish peroxidase-conjugated (HRP) α -mouse IgG and HRP α -rabbit IgG from Vector Laboratories (Burlingame, CA, USA); FITC-conjugated goat α -mouse IgG (Thermo Scientific); NMS-873, ML-240 and DBeq from Cayman Chemicals (Ann Arbor, MI, USA); propidium iodide (PI) from Santa Cruz Biotechnology (Dallas, TX, USA) and α -VCP 612182 from BD Biosciences (Franklin Lakes, NJ, USA). α - β -tubulin E7 was obtained from Developmental Studies Hybridoma Bank (DBHS, University of Iowa, USA). NMS-873, ML-240, DBeq and progesterone were dissolved in DMSO; PI was dissolved in hexa-distilled water.

Animals

Hybrid F1 (C57BL/6 male x BALB/c female) mature (8–12 weeks-old) mice, as well as mature transgenic [BDF1-Tg (CAG-mtDsRed2, Acr3-EGFP) RBGS0020sb] male mice who have sperm with acrosomal vesicles expressing green EGFP fluorescence and mitochondria expressing red DsRed2 fluorescence (Hasuwa et al., 2010), were used. Animals were maintained at 23°C with a 12 hr light: 12 hr dark cycle. Animal experimental procedures were reviewed and approved by the Ethical Committee of IBYME (CE/003-1/2011). Experiments were performed in strict accordance with the Guide for Care and Use of Laboratory Animals approved by the National Institutes of Health (NIH).

Sperm capacitation

The non-capacitating medium used in this study was a modified TYH medium, containing 119.3 mM NaCl, 4.7 mM KCl, 1.71 mM $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$, 1.2 mM KH_2PO_4 , 1.2 mM $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$, 0.51 mM sodium pyruvate, 5.56 mM glucose, 20 mM 4-(2-hydroxyethyl) piperazine-1-ethanesulfonic acid (HEPES) and 10 $\mu\text{g}/\text{ml}$ gentamicin (NC medium). This medium, which does not support capacitation, was prepared without BSA and NaHCO_3 . Animals were euthanized and cauda epididymal mouse sperm were collected and placed in NC medium. After 15 min of incubation at 37°C, epididymides were removed, and sperm were resuspended to a final maximum concentration of 10^7 cells/ml in the appropriate medium depending on the experiment performed. For capacitation, a medium containing 5 mg/ml of BSA and 15 mM of NaHCO_3 was used, and sperm were incubated at 37°C (CAP medium) for 90 min. In the cases where medium without added Ca^{2+} salts were required, 1.71 mM $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$ was omitted (nominal zero- Ca^{2+} medium, indicated by $\text{Ca}^{2+} = 0$). When chelation of the extracellular Ca^{2+} was needed, 1 mM EGTA was added at nominal

zero- Ca^{2+} medium (indicated by $\text{Ca}^{2+}=0 + \text{EGTA}$). In all cases, pH was adjusted to 7.4 with NaOH. To test the effect of inhibitors on capacitation, sperm were pre-incubated with inhibitors in the NC medium for 10 min prior to the beginning of the capacitating period.

Immunoblotting

After incubation in the appropriate medium, sperm were washed by centrifugation (5 min at 400 g), resuspended in sample buffer without reducing agents (62.5 mM Tris HCl pH 6.8, 2% SDS, 10% glycerol), and boiled for 5 min. After centrifugation for 5 min at 13,000 g, 5% β -mercaptoethanol and 0.0005% bromophenol blue were added to the supernatants then boiled again for 5 min. Protein extracts equivalent to $2\text{--}4 \times 10^6$ sperm per lane were separated by SDS-PAGE in gels containing 10% polyacrylamide and transferred onto nitrocellulose membranes. Blots were blocked in 5% nonfat dry milk in PBS containing 0.1% Tween 20 (T-PBS) for 1 hr at room temperature. An overnight incubation at 4°C with the first antibody was required. Antibodies were diluted in 2% nonfat dry milk in T-PBS as follows: α -VCP (1:1000), α -pPKAs (1:3000), α -pY (1:3000) and α - β -tubulin (1:3000). The corresponding secondary antibodies were incubated for 45 min at room temperature, diluted in 2% nonfat dry milk in T-PBS as follows: 1:1000 for HRP α -rabbit and 1:4000 for HRP α -mouse. In all the cases, the reactive bands were detected by enhanced chemiluminescence (GE Healthcare, Piscataway, NJ, USA). When necessary, membranes were stripped for 10 min in 2% SDS, 0.67% β -mercaptoethanol, 62.5 mM Tris pH 6.8. In all experiments, molecular masses were expressed in kilodaltons (kDa). As a loading control, α - β -tubulin was used. For band quantification, rectangular boxes were drawn around bands on scanned digital images of ECL contact photographs of Western blots. Adjusted optical densities for each lane were obtained using ImageJ (NIH, USA).

Immunoprecipitation

Immunoprecipitation was performed as previously described with minor modifications (Stival et al., 2015). After sperm incubation in the appropriate conditions, samples were centrifuged at 1,700 g for 1 min. The resulting pellet was resuspended in RIPA buffer (10 mM Tris-HCl pH 7.2, 50 mM NaCl, 0.1% SDS, 1% Triton X-100, 1 mM EDTA, 1 mM sodium orthovanadate and protease inhibitors), incubated on ice for 30 min and centrifuged at 4°C for 5 min at 2,500 g. Supernatants were incubated sequentially (after stripping) with α -VCP or α -pY antibodies (1:200 of antibody for 10^7 cells in a final volume of 500 μ l) for 2 hr at room temperature with constant rocking. After adding 20 μ l of protein G-sepharose (GE Healthcare), the reactions were further rocked for 1 hr at 4°C. The immune complex was recovered by centrifugation, washed four times in RIPA buffer, and subjected to SDS/PAGE and Western blot. To avoid immuno-reactive signals from denatured IgGs used for immunoprecipitation, a monoclonal secondary-HRP mouse α -native rabbit IgG was used (clone RabT-50, Sigma). Input refers to the total protein used for IP.

Immunofluorescence and super-resolution microscopy

To perform immunofluorescence on mouse sperm, we used a previously described method (Romarowski et al., 2015). Briefly, mouse sperm obtained from cauda epididymis and capacitated during different times were washed twice by centrifugation for 5 min at 400 g, resuspended in PBS, and placed onto cover slides. Sperm were air-dried and then treated

with PBS containing 4% formaldehyde for 15 min at room temperature. After washing twice for 5 min in T-PBS, sperm were permeabilized with 0.5% Triton X-100 in PBS for 5 min. The slides were then washed twice for 5 min and incubated in T-PBS containing 10% normal goat serum (blocking buffer) for 1 hr at room temperature, then with primary antibody (1:100; purified mouse α -VCP) diluted in T-PBS containing 1% normal goat serum for an overnight period at 4°C. Following a washing step (three times, 5 min in T-PBS), slides were incubated with FITC-conjugated goat α -mouse IgG diluted 1:200 in T-PBS containing 1% normal goat serum for 1 h at room temperature, washed again for three times, and mounted using Vectashield (Vector Labs, Burlingame, CA, USA). Non-specific staining was determined by incubating the sperm with rabbit IgG. Samples were examined using either a laser scanning confocal microscope (Nikon C1 Eclipse 800) equipped with a 60X objective (NA 1.4) or a NanoImager S microscope (Oxford Nanoimaging Ltd), equipped with a 100X oil-immersion objective (NA 1.4, Olympus) and a sCMOS camera (Hamamatsu ORCA-Flash4.0 V3 Digital CMOS camera).

CatSper opening through determination of membrane potential (E_m) by flow cytometry

Sperm E_m changes were assessed using DiSC₃(5) as previously described (Luque et al., 2023; Puga Molina et al., 2018). Briefly, after a 20-min incubation with VCP inhibitors (20 μ M NMS-873, 1 μ M ML-240 or 5 μ M DBeq), CatSper inhibitor (as a negative control, 20 μ M HC-056456) or vehicle (DMSO), 50 nM DiSC₃(5) was added. In these experiments, DiSC₃(5) was not washed. To monitor viability, 3 μ M of PI was added before measurement. Data were recorded as individual cellular events using a BD FACSCanto II™ cytometer (Biosciences; Becton, Dickinson and Company). Basal E_m was obtained after 30 s of continuous recording, followed by chelation of Ca²⁺ with 3.5 mM of EGTA. Then, acquisition continued for an additional 150 s. Positive cells for DiSC₃(5) were collected using the filter for allophycocyanin (APC; 660/20), and for PI using the filter for peridinin chlorophyll protein complex (PerCP; 670LP). Compensation was done. Data were analyzed using FlowJo software (V10.0.7).

Determination of intracellular cAMP levels

After the incubation of sperm in the appropriate conditions for 12 min, reactions were stopped by centrifugation at 2,000 *g* for 3 min at room temperature. Cells were resuspended with 200 μ l of 0.1 M HCl and lysed for 10 min at room temperature. Sperm lysates were centrifuged at 2000 *g* for 3 min and the cAMP in the supernatant was acetylated and quantified using the Direct cAMP ELISA Kit (Enzo, Farmingdale, NY, USA; Catalog # ADI-901-066A), according to the manufacturer's instructions. This ELISA-based assay has an increased sensitivity of 0.006 pmol/ml cAMP in the acetylated assay format. A standard curve was run for each assay and the unknown cAMP concentrations were obtained by interpolation as recommended by the manufacturer.

Computer-assisted sperm analysis (CASA)

After incubation in the appropriate conditions, sperm suspensions were loaded on a 100- μ m depth chamber slide (Leja Slide, Spectrum Technologies) and placed on a microscope (Nikon Eclipse E200) stage at 37°C coupled to a Basler acA780-75gc camera. Sperm movements were examined using a CASA system (Sperm Class Analyzer: SCA evolution,

Microptic, Barcelona, Spain). Parameters used were as follows: 30 frames acquired, frame rate of 60 Hz, and cell size of 30-170 μm^2 . At least 20 microscopy fields corresponding to a minimum of 200 sperm were analyzed in each experiment. The following parameters were measured: mean path velocity (VAP), curvilinear velocity (VCL), straight line velocity (VSL), linearity (LIN), amplitude of lateral head displacement (ALH), and straightness (STR).

Acrosomal exocytosis

Acrosome reaction was assessed as previously described (Hirohashi et al., 2015). Briefly, mt-DsRed2 Acr-EGFP mouse sperm were resuspended to a final maximum concentration of 10^7 cells/ml on NC medium. A 10-min preincubation in NC medium containing inhibitors was conducted when required. After that, sperm were incubated under CAP conditions for 60 min in the presence or absence of inhibitors. At 60 min of incubation, progesterone (30 μM) or A23187 (10 μM) was added for 30 min to induce the acrosome reaction. Before collecting data, 2 $\mu\text{g}/\text{ml}$ of propidium iodide was added to monitor viability. Data were recorded as individual cellular events using a BD FACSCanto II TM cytometer (Biosciences; Becton, Dickinson and Company). Side-scatter area and forward-scatter area data were collected from 20,000 events per sample to define sperm population. In all cases, doublet exclusion was performed by analyzing the two-dimensional dot plot forward-scatter area versus forward-scatter height. Positive cells for EGFP were collected using the filter for FITC (530/30) and for propidium iodide, using the filter for peridinin chlorophyll protein complex (670LP). The two indicators had minimal emission overlap, but compensation was still done. Data were analyzed using FlowJo software (V10.0.7).

In vitro fertilization (IVF)

Eight to 12-week-old F1 female mice were superovulated using equine chorionic gonadotropin (5 IU, PMSG; Syntex), followed 48 hr later by human chorionic gonadotropin (5 IU, hCG; OVUSYN, Syntex) intraperitoneal injection. Cumulus-oocyte complexes (COCs) were collected from oviducts 13-14 hr after hCG administration, placed in TYH IVF medium (containing 25 mM NaHCO_3 and 4 mg/ml of BSA, with no HEPES addition). Sperm were capacitated *in vitro* at a final concentration of 5×10^5 sperm/ml in the presence or absence of 20 μM NMS-873. Once capacitated, sperm were washed by centrifugation for 5 min at 400 g, resuspended in fresh TYH CAP medium, and used to inseminate COCs. After coincubation for 4 hr at 37°C with 5% CO_2 , eggs were washed three times and placed in drops containing TYH IVF at 37°C with 5% CO_2 . Fertilization was assessed by the visualization of 2-cell embryos 20 hr later.

Statistical analysis

Data are expressed as mean \pm standard error of the mean (SEM) of at least three experiments for all determinations. Statistical analyses were conducted using GraphPad Prism v4.0 (San Diego, CA, USA). The specific statistical analysis employed is indicated in the figure legends. A p-value of less than 0.05 ($p < 0.05$) was considered statistically significant.

Results

VCP localized to the equatorial segment and to the flagellum of mouse sperm

In mouse sperm, the α -VCP antibody recognized a single band for the expected size (97 kDa) by Western blot (Figure 1A). Immunolocalization revealed remarkable differences compared to human sperm (Ficarro et al., 2003). VCP is localized in the equatorial segment of the mouse sperm head and in the sperm flagellum (Figure 1B). Localization in the head was confirmed using transgenic Acr3-EGFP-sperm, with the strong VCP signal in the equatorial segment while EGFP was mostly restricted to the anterior acrosomal cap (Figure 1C).

VCP is not tyrosine phosphorylated during mouse sperm capacitation

In contrast to what was reported in human sperm (Ficarro et al., 2003), mouse sperm VCP did not change its equatorial segment localization during capacitation (Figure 2A). Because of the differences observed in the localization of VCP in mouse and human sperm, we investigated if VCP is tyrosine phosphorylated during capacitation. To that end, an α -VCP was used for immunoprecipitation of non-capacitated as well as of capacitated sperm protein extracts. Sperm of each population were extracted and immunoprecipitated using α -VCP and the immunoprecipitates were then probed with both α -pY or α -VCP by immunoblotting, as shown in Figure 2B. Overall, this experiment confirmed that VCP was effectively immunoprecipitated by the α -VCP antibody since a single band of the expected molecular size (97 kDa) was detected in both non-capacitated (NC) and capacitated (CAP) sperm extracts (Figure 2B, left panel). However, these bands were not observed when the membrane was probed with an α -pY antibody (Figure 2B, right panel) indicating that VCP does not undergo tyrosine phosphorylation during capacitation.

VCP is released and cleaved during the course of the acrosome reaction

Due to the localization of VCP in the equatorial segment of mouse sperm, we investigated its fate during either induced or spontaneous acrosome reaction. To this end, transgenic Acr-EGFP exposed to agonists of acrosome reaction (progesterone or the Ca^{2+} ionophore A23187) were fixed and stained with α -VCP. In Figure 3A, a representative set of images corresponding to this experiment is shown. Most of the acrosome-intact sperm exhibited VCP in the equatorial segment. However, those that underwent acrosome reaction lost the VCP signal in the equatorial segment (see quantification in Figure 3B). Despite VCP being lost after acrosome reaction, the signal in the flagellum remained unaltered, as shown in Figure 3C.

Next, we investigated if this protein is released to the surrounding medium. After the incubation in capacitating conditions, the suspension of sperm undergoing spontaneous or induced acrosome reaction was centrifuged, and the supernatant was analyzed by immunoblotting using α -VCP. We observed the presence of a 58 kDa VCP fragment in the supernatant of sperm incubated under capacitating conditions in those that underwent either induced or spontaneous acrosome reaction, suggesting the release of this fragment (Figure 3D). In contrast, the presence of the full-length 97 kDa form was not observed. A previously reported *in silico* analysis of the VCP sequence demonstrated the presence of

2 cleavage sites for trypsin-like proteases that produce VCP fragments of 58 and 87 kDa (Halawani et al., 2009), which match those observed in our experimental conditions (Figure 3D). As a control, media recovered from sperm that swam out from the cauda epididymis in non-capacitating medium did not show the presence of VCP evidenced by the absence of the band corresponding to VCP. The 87-kDa form is barely observed in those sperm released from the cauda epididymis (see first lane) and could not be observed after exposure to capacitating medium.

Inhibition of VCP led to reduced protein tyrosine phosphorylation

We further explored the role of VCP in the flagellum using different pharmacological inhibitors against this protein. These inhibitors possess different mechanisms of action. NMS-873 is a highly selective allosteric inhibitor that acts by binding to the interface of two adjacent domains within the active hexameric structure of VCP and leads to an interruption of its catalytic cycle by stabilizing the ADP-bound state (Bouwer et al., 2021). As for DBeq, it was identified as a selective, potent, reversible, and ATP-competitive VCP inhibitor that targets both D1 and D2-ATPase domains of VCP (Chou et al., 2011; Fang et al., 2015). In contrast, ML-240, derived from the scaffold of DBeq, only targets the D2-ATPase domain (Fang et al., 2015).

We used these inhibitors to investigate if VCP plays a role in regulating phosphorylation events that take place in the flagellum mediated by the cAMP/PKA pathways. As depicted in Figure 4A and Supplementary Figure S1, the incubation of sperm for 90 min under capacitating conditions with increasing concentrations of the three inhibitors resulted in reduced tyrosine phosphorylation levels in a concentration-dependent manner. The maximum concentration tested for each VCP pharmacological inhibitor was chosen as 20 μM for NMS-873, 1 μM for ML-240, and 5 μM for DBeq.

To better understand the role of VCP in PKA activity, we subjected sperm to a challenge by chelating extracellular Ca^{2+} . In the complete absence of extracellular Ca^{2+} , as seen in conditions with EGTA, tyrosine phosphorylation levels experience a significant increase, but only if PKA is active (Navarrete et al., 2015). Consequently, in Figure 4B and Supplementary Figure S2, the presence of VCP inhibitors leads to a decrease in tyrosine phosphorylation when exposed to EGTA, providing further support for the notion that VCP operates upstream of PKA action.

Partial inhibition of CatSper channels by VCP inhibitors might lead to a decrease in tyrosine phosphorylation, akin to the conditions observed when sperm are incubated with low concentrations of Ca^{2+} (Navarrete et al., 2015). One way of evaluating CatSper activity indirectly is by measuring E_m changes after removing external free divalent cations by chelation with EGTA. This will cause CatSper to efficiently conduct monovalent cations such as Na^+ when it is open. Since Na^+ is highly concentrated in the extracellular milieu, a sudden influx depolarizes the cell, and it can be measured with the cationic dye DiSC₃(5) by flow cytometry. Our experiments depicted in Supplementary Figure S4 A demonstrate that VCP inhibitors do not impact CatSper function, evidenced by the lack of alteration in the depolarization of the E_m in response to EGTA (Luque et al., 2021; Luque et al., 2023), in contrast to the CatSper inhibitor HC-056456.

Diminished cAMP levels and PKA activation as a result of VCP inhibition

To investigate if VCP inhibition alters the capacitation-induced PKA activation, sperm were incubated under capacitating conditions in the presence of the VCP inhibitor NMS-873, and PKA-dependent phosphorylation was evaluated. Incubation with NMS-873 led to a decrease in the phosphorylation of PKA substrates (Figure 5A) suggesting that active VCP is essential for proper PKA activity. To verify that the action of VCP is upstream PKA activation, sperm were incubated under capacitating conditions with NMS-873, followed by the addition of increasing concentrations of the membrane-permeable cAMP analog, 8-Br-cAMP. Under these conditions, the levels of PKA substrates and tyrosine phosphorylation were increased in a concentration-dependent manner reaching levels comparable to the control capacitating condition (Figure 5A). The same behavior of a decrease in PKA substrates phosphorylation is seen when Dbeq and ML-240 are used to inhibit VCP and, as observed with NMS-873, this inhibition is rescued by increasing concentrations of 8-Br-cAMP (Supplementary Figure S4 B). These findings indicate that VCP regulates the signals essential for the occurrence of PKA activation, acting upstream PKA.

To further explore the role of VCP on PKA activation, intracellular cAMP levels were examined when sperm were exposed to both non-capacitating and capacitating conditions, while being treated with NMS-873. As illustrated in Figure 5B, intracellular cAMP levels exhibited a rapid increase when sperm were incubated under capacitating conditions. However, the presence of the VCP inhibitor NMS-873 entirely abolished this increase in intracellular cAMP, resulting in levels akin to those observed in non-capacitating conditions. Altogether, these results suggest that VCP activity is required for maintaining the capacitation-associated rise in intracellular cAMP levels.

VCP inhibition impairs the rate of acrosomal exocytosis and decreases the fertilization rate in IVF without affecting sperm motility

Sperm were subjected to incubation under capacitating conditions while exposed to increasing concentrations of the VCP inhibitor NMS-873. As illustrated in Supplementary Figure S4 C and Table S1, there were no significant differences observed in sperm motility as assessed by computer-assisted sperm analysis (CASA).

Acrosome reaction was assessed by exposing sperm to increasing concentrations of NMS-873 when incubated under capacitating conditions followed by addition of either 30 μ M progesterone or 10 μ M A23187. As shown in Figure 6A, a progressive inhibition of the progesterone-induced acrosome reaction was observed, and it became statistically significant at the concentration of 2 μ M. In contrast, no inhibition was noted when sperm were challenged with calcium ionophore.

Finally, IVF experiments were conducted in the presence of 20 μ M NMS-873, and the results were compared to those of the control group (vehicle). As shown in Figure 6B, sperm that had been pre-incubated with the VCP inhibitor exhibited lower fertilization rates when compared to the control sperm.

Discussion

In the present work, we analyzed the role of VCP in mouse sperm. First, we found that the localization of this protein in different compartments of the mature sperm. The first evidence related to this protein in mammalian sperm shows that it is localized in the neck and tail of human sperm, and that during capacitation, VCP is relocalized to the anterior acrosome (Ficarro et al., 2003). In mouse sperm, we observed a different localization; in this species VCP localized to the equatorial segment and the flagellum, and this pattern is not altered during capacitation. The area corresponding to the equatorial segment of the sperm head is of great importance for mammalian fertilization since fusion between the sperm plasma membrane and the egg's oolema takes place in this region. As such, key proteins for sperm-egg fusion such as IZUMO1 are relocalized to this region after acrosome reaction (Miranda et al., 2009; Satouh et al., 2012). Other proteins that have been reported to migrate to the equatorial segment in mouse sperm are CD81 (Jankovicova et al., 2016), CAPZA3 (Sosnik et al., 2010), CRISP1/2 (Busso et al., 2007), α -L-Fucosidase, SLLP1 (Herrero et al., 2005; Mandal et al., 2003), fertilin β (Hunnicutt et al., 1997), and equatorin (Toshimori et al., 1992). However, to the best of our knowledge, with the exception of TMEM225 (Matsuura and Yogo, 2015) and Capza3 (Sosnik et al., 2010), all the proteins that are a constitutive part of the equatorial segment or that are relocalized to this region are not lost after the acrosome reaction. In fact, it is possible that the major reorganization that takes place in the plasma membrane during capacitation promotes the breakdown of the diffusion barriers between domains, allowing protein migration to the equatorial segment (Flesch FM et al., 2000; Gadella, 2008). Proteins located in the acrosome, on the other hand, could either migrate to the equatorial segment via the "hairpin" structure formed by the acrosomal and plasma membranes in the equatorial sheath after the acrosome reaction (Leung et al., 2021). Here, we show that VCP is lost after the acrosome reaction and further experiments are needed to clarify in detail the subcellular localization and understand the mechanisms by which some proteins are lost while others remain attached to this area.

As previously mentioned, in humans VCP is phosphorylated in tyrosine residues during capacitation (Ficarro et al., 2003). In this manuscript, it was speculated that this phosphorylation may be linked to the migration of VCP to the acrosomal cap. This hypothesis is based on results obtained in other systems where VCP undergoes tyrosine phosphorylation during T-cell activation. In these cells, although phosphorylation did not alter its ATPase activity, it regulates the subcellular localization of this protein. In the mouse, VCP is not phosphorylated in tyrosine residues during capacitation, which may be consistent with the lack of change in localization during capacitation.

Another finding that emerged from our experiments is that VCP is cleaved after it is released to the surrounding medium. A previously published *in silico* analysis suggests that the observed fragments are compatible with target sequences for trypsin or trypsin-like proteases (Halawani et al., 2009). One of the sperm-specific trypsin-like protease is acrosin. This enzyme is one of the many acrosomal proteases responsible for digesting specific acrosomal components such as ZP3R/sp56 (Buffone et al., 2008; Buffone et al., 2009).

Regarding the role of VCP in the sperm flagellum, where most of the tyrosine phosphorylated and PKA-induced phosphorylated substrates localized, experiments using inhibitors against VCP showed that PKA substrates and tyrosine phosphorylation are substantially reduced. In the complete absence of Ca^{2+} such as in CatSper KO sperm, in the presence of CatSper inhibitors or by adding EGTA, tyrosine phosphorylation levels increase dramatically. Our hypothesis is that VCP is positively helping the occurrence of tyrosine phosphorylation. To further support this hypothesis, extracellular Ca^{2+} was removed by addition of EGTA. This treatment greatly increases the levels of tyrosine phosphorylation when PKA is active (Navarrete et al., 2015). However, in conditions where VCP was inhibited, the EGTA-induced increase was almost completely obliterated.

Capacitation, and therefore, the preparation for the acrosome reaction is also regulated by cAMP-dependent pathways (Visconti et al., 1995). Thus, the altered induction of the acrosome reaction is consistent with the low levels of cAMP assessed in the presence of VCP inhibitors. Similarly, the inhibition of PKA-dependent phosphorylation explains the block on tyrosine phosphorylation, a process mediated by FERT (Alvau et al., 2016). Also, the block of capacitation-induced cAMP-dependent pathway is consistent with the significant reduction of IVF rates. We did not observe a complete block of fertilization in these conditions; this result is probably due to dilution of the inhibitor concentration in the fertilization droplet.

In conclusion, our results are consistent with VCP involvement in the regulation of cAMP-dependent pathways. In addition, VCP cleavage and loss from the equatorial segment after the acrosome reaction suggests alternative regulation of this protein during exocytosis. Future experiments will be necessary to unravel the molecular mechanisms underlying VCP's involvement in these two functions.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgements

We would like to thank Rene Baron and Williams Foundation. We thank Adan Guerrero and Alberto Darszon for their insightful comments.

Disclosure of Interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported. Mariano G Buffone is an Associate Editor] of *Reproduction*. Mariano G Buffone was not involved in the review or editorial process for this paper, on which he is listed as an author

This work was supported by the National Institutes of Health (NIH R01HD106968 to MGB and DK; NIH-HD088571 to PEV and HD38082 to PEV), *Agencia Nacional de Promoción Científica y Tecnológica* (PICT 2020-00098, 2021-00031) *Consejo Nacional de Investigaciones Científicas y Técnicas* (PIBAA 2022-100368 to GML) and Chan Zuckerberg Initiative (CZI 2021–240504).

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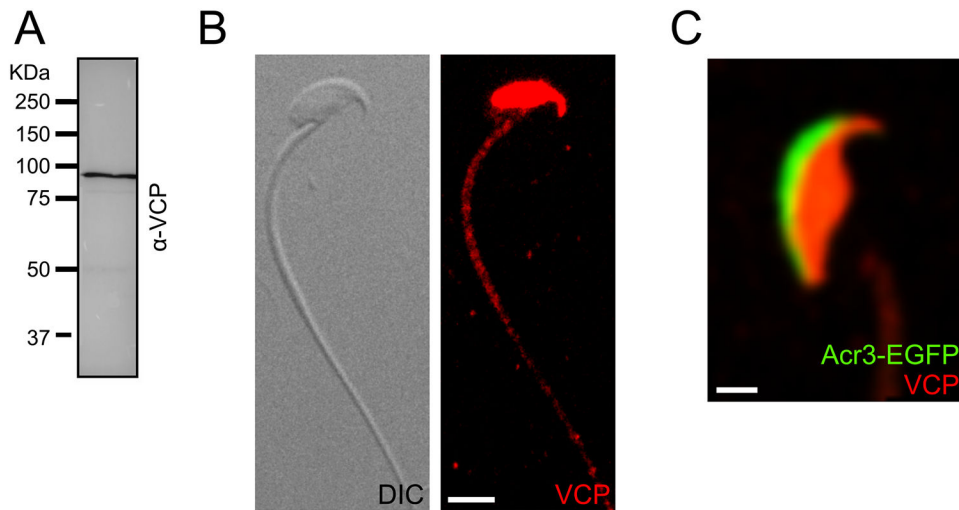


Figure 1: VCP is expressed in mouse sperm and localized in the equatorial segment and the flagellum.

A) Mouse sperm proteins were analyzed by 10% SDS-PAGE and immunoblotted using an antibody against VCP. The lane contained 5×10^6 sperm. **B)** Immunofluorescence showing the localization of VCP in mouse sperm incubated under non-capacitating (NC) conditions. Scale bar = 5 μm . A representative image of at least 3 independent experiments is shown. **C)** Immunofluorescence showing the localization of VCP in transgenic Acr3-EGFP mouse sperm. Scale bar = 2 μm .

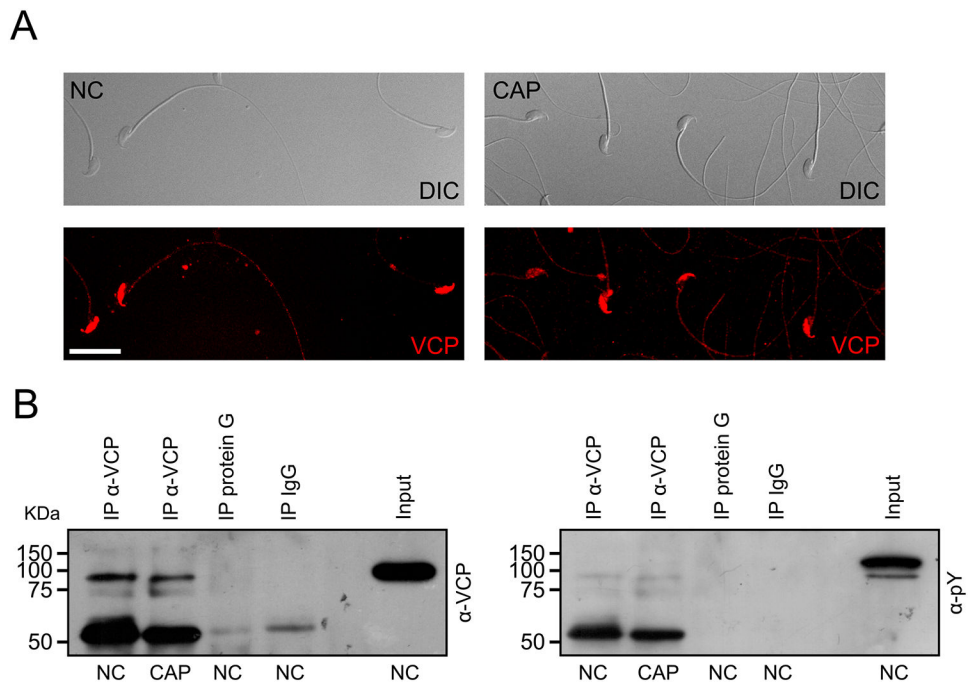


Figure 2: VCP is not phosphorylated in tyrosine residues during capacitation.

A) Immunofluorescence showing the localization of VCP in mouse sperm incubated under capacitating (CAP) or non-capacitating (NC) conditions. Representative images of at least 3 independent experiments are shown. Scale bar = 20 μ m. **B)** Sperm incubated in capacitating and non-capacitating media were extracted with RIPA buffer and immunoprecipitated with α -VCP. Immunocomplexes were subjected to 10% SDS/PAGE, transferred to PVDF membranes and developed two times (after stripping) with either α -VCP (left panel) or α -pY (right panel). Input refers to the total protein used for IP.

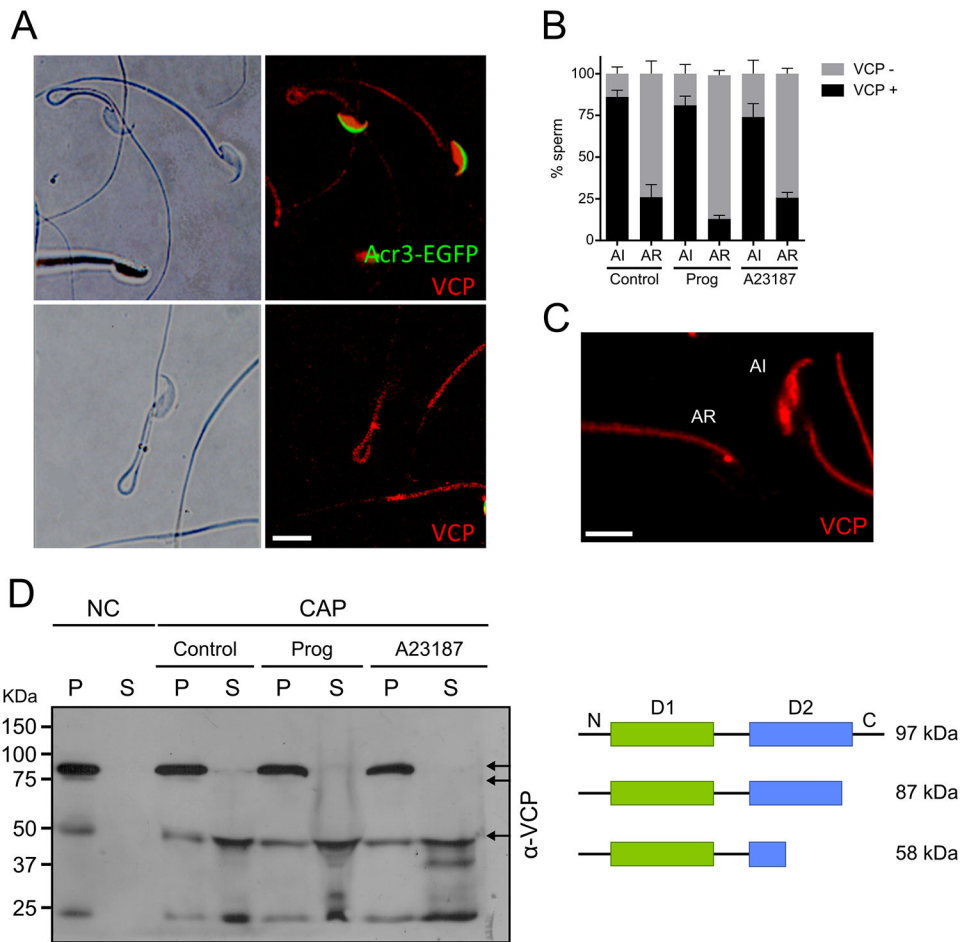


Figure 3: VCP is cleaved and released from the ES during AR.

A) Capacitated Acr3-EGFP sperm were incubated under CAP conditions followed by exposure to 30 μ M progesterone to induce acrosome-reaction. Then, sperm were fixed, attached to a slide and stained with α -VCP antibody followed by α -mouse TRITC. Slides were observed using a confocal microscope. Scale bar = 10 μ m. **B)** Percentage of sperm that display a positive or negative signal for VCP (VCP+, VCP-, respectively) in the equatorial segment in the acrosome-intact (AI) or acrosome-reacted (AR) population. Sperm were analyzed in different conditions: after the incubation under CAP conditions (Control); after the incubation under CAP conditions and stimulated with either progesterone (Prog 30 μ M) or A23187 (10 μ M). Data from at least 3 independent experiments are shown, with 100 cells analyzed. **C)** Immunofluorescence showing the localization of VCP in AI and acrosome-reacted (AR) mouse sperm. Note how, although lost in the head, VCP remains in the flagellum after acrosome-reaction. Scale bar = 5 μ m. **D) Left panel: Equal number of mouse sperm** were incubated under NC and CAP conditions for up to 90 min. A subset of cells was then stimulated with progesterone (30 μ M) or A23187 (10 μ M). Then, the incubation mixture was separated into a pellet (P) and supernatant (S) and analyzed by 10% SDS-PAGE and immunoblotting with α -VCP. The α -VCP antibody detected molecular bands of ~97, ~87 and 58 kDa. The experiment was repeated at least 3 times. **Right panel:**

Schematic representation of VCP in mouse sperm. The 97 kDa monomer is composed of 2 contiguous domains (domain 1 D1 and domain 2 D2)(Halawani et al., 2009).

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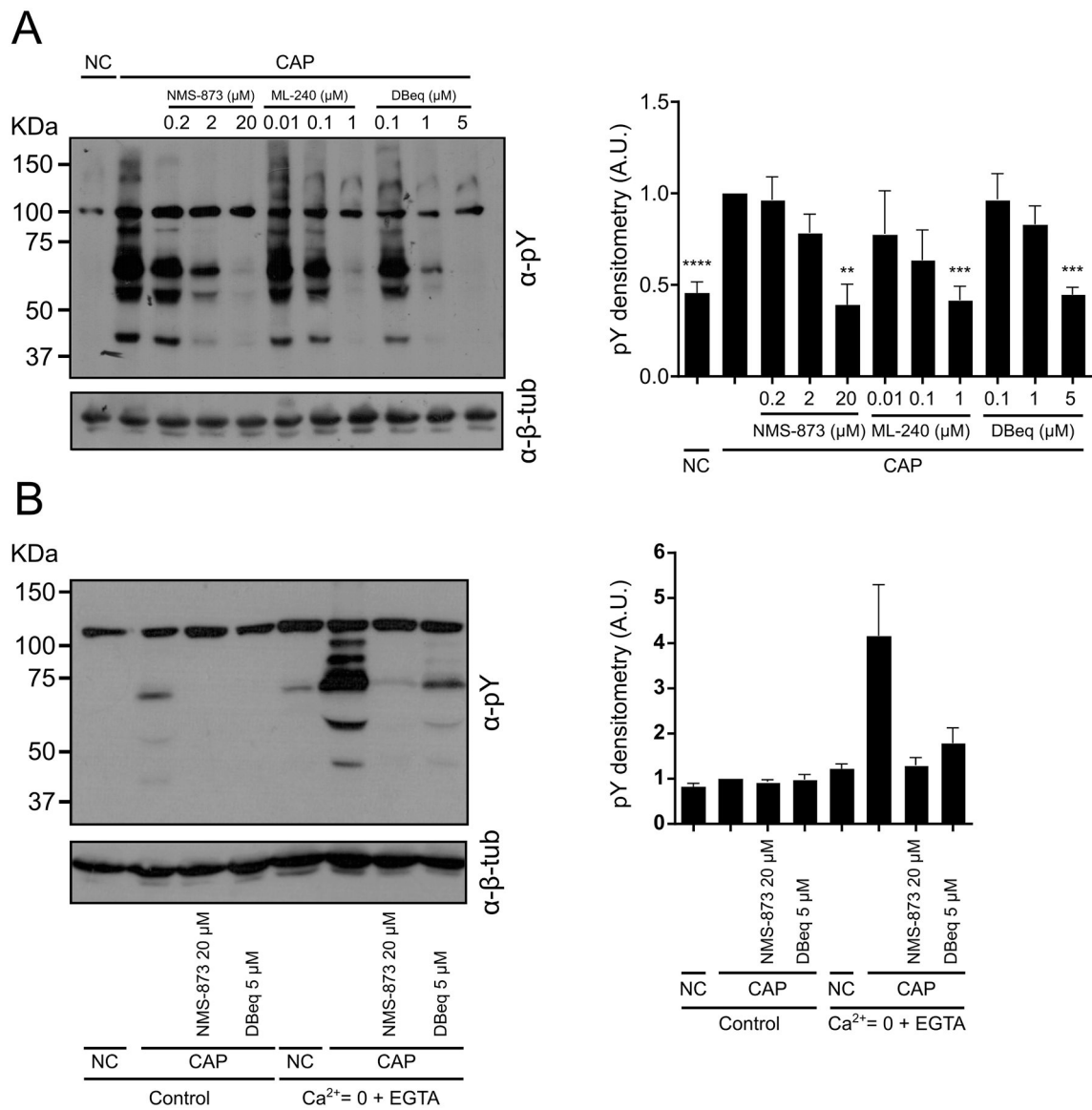


Figure 4. VCP inhibition abolishes sperm protein tyrosine phosphorylation during capacitation
A) Protein extracts were separated by 10% SDS-PAGE and immunoblotted with an antibody against pY. As a loading control, β-tubulin was used. Sperm were incubated for 90 min under NC or CAP conditions in the absence (DMSO) or presence of VCP inhibitors. Quantitative analysis was performed by measuring the optical density of all bands and relativized to β-tubulin. Data is presented as the mean ± SEM of at least 3 independent experiments, where normalization to the control condition (CAP 90 min with DMSO) was used. ****p<0.0001 represents statistical significance vs control (CAP 90 min with DMSO). Non-parametric Kruskal-Wallis test was performed in combination with Dunn's multiple comparisons test. **B)** Protein extracts were separated by 10% SDS-PAGE and immunoblotted with an antibody against pY. As a loading control, β-tubulin was used. Sperm were incubated for 90 min under NC and CAP in the absence (DMSO) or presence of VCP inhibitors (20 μM NMS-873 and 5 μM DBeq). In addition, two different media were

used: a) complete TYH medium (Control); b) medium without added Ca^{2+} salts (nominal zero- Ca^{2+} medium, $\text{Ca}^{2+}=0$) with EGTA 1 mM ($\text{Ca}^{2+}=0$ + EGTA). Quantitative analysis was performed by measuring the optical density of all bands and relativized to β -tubulin. Data is presented as the mean \pm SEM of at least 3 independent experiments, where normalization to the control condition (CAP 90 min with DMSO) was used. Non-parametric Friedman test was performed in combination with Dunn's multiple comparisons test.

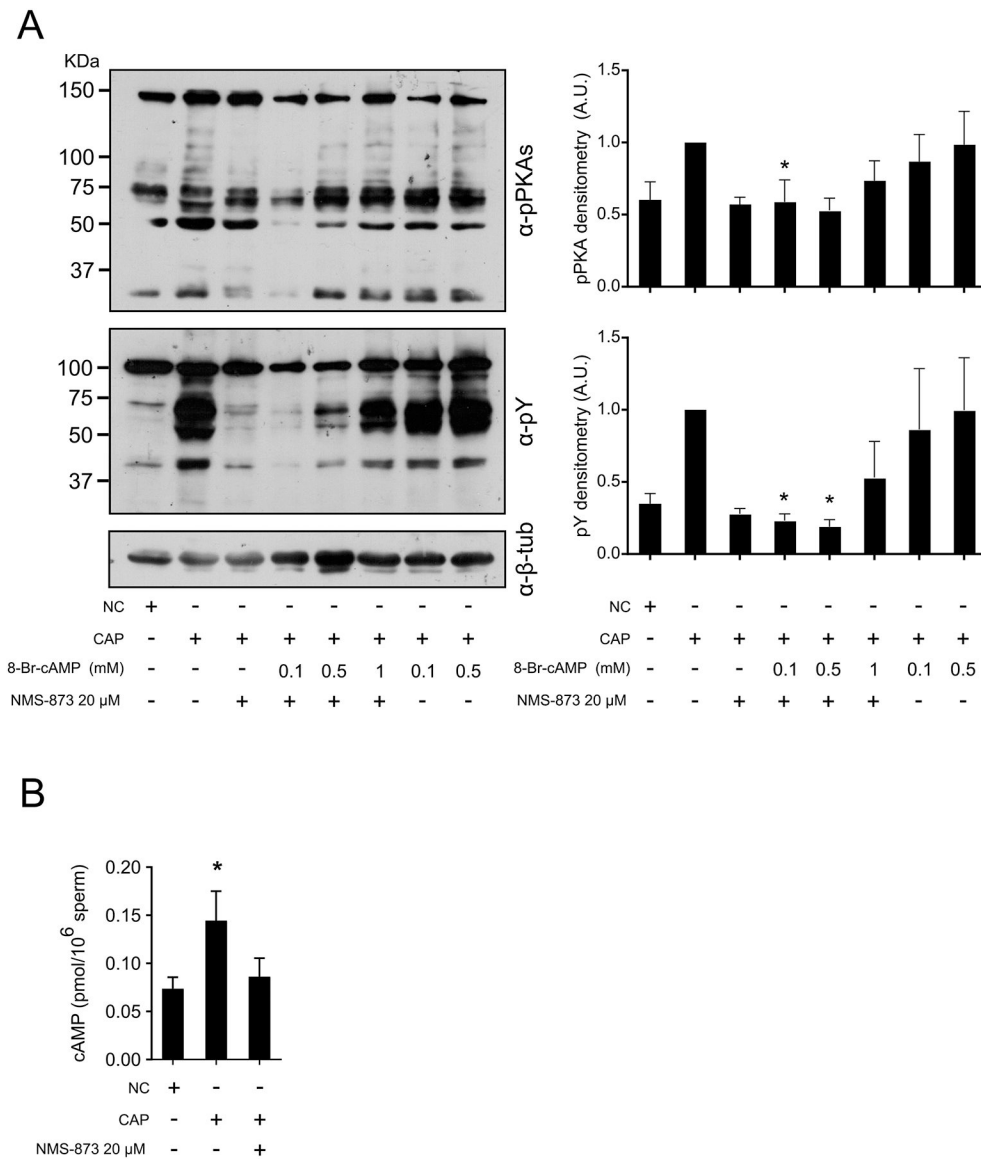


Figure 5. cAMP levels are reduced in the presence of VCP inhibitors.
A) Protein extracts were separated by 10% SDS-PAGE and immunoblotted with antibodies against substrates phosphorylated by PKA (pPKAs) and pY. As a loading control, β -tubulin was used. Sperm were incubated for 90 min under NC or CAP in the absence (DMSO) or presence of 20 μ M NMS-873 together with increasing concentrations of 8-Br-cAMP (0.1; 0.5 and 1 mM). Quantitative analysis was performed by measuring the optical density of all bands and relativized to β -tubulin. Values represent the mean \pm SEM of at least 3 independent experiments, where normalization to the control condition (CAP 90 min with DMSO) was used. * $p < 0.05$ represents statistical significance vs control (CAP 90 min with DMSO). Non-parametric Friedman test was performed in combination with Dunn's multiple comparisons test. **B)** Total intracellular cAMP content of sperm incubated for 90 min under NC or CAP conditions with or without 20 μ M NMS-873. Values represent the mean \pm SEM of 4 independent experiments. * $p < 0.05$ represents statistical significance vs control (CAP 90 min with DMSO).

min with DMSO). Non-parametric Friedman test was performed in combination with Dunn's multiple comparisons test.

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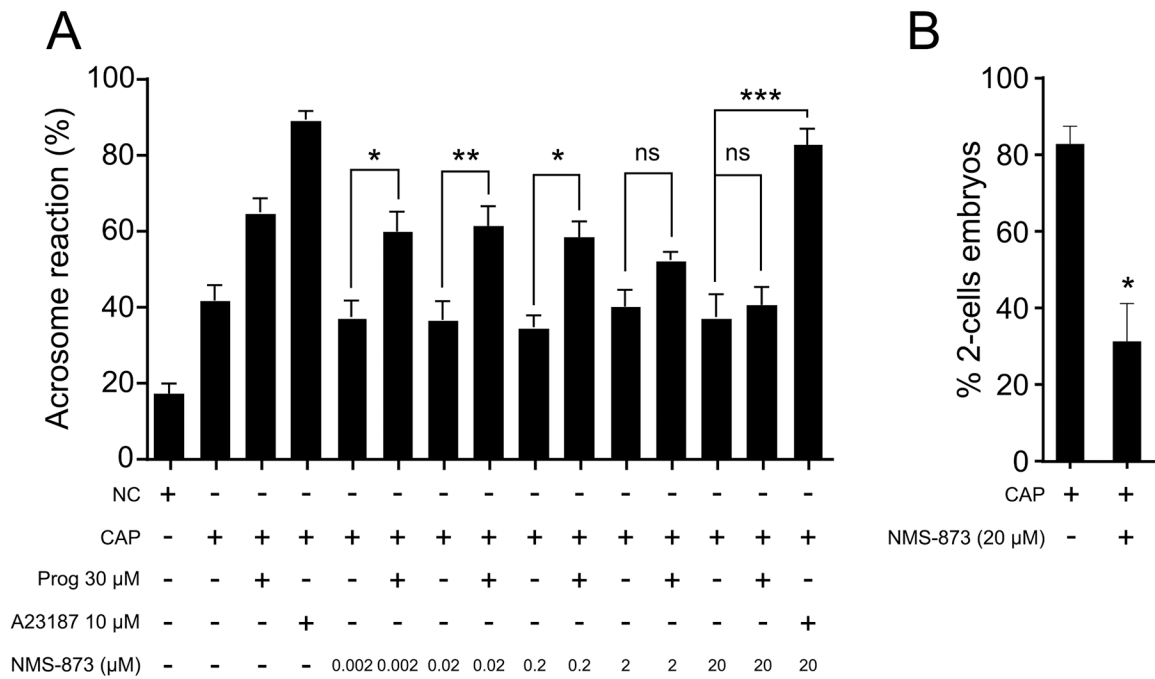


Figure 6. Effect of VCP inhibitor NMS-873 on acrosome reaction and IVF.

A) Acrosome reaction was assessed by flow cytometry analysis of Acr-EGFP sperm. Mouse sperm were incubated for 90 min under capacitating conditions in the absence (DMSO) or presence of increasing concentrations of NMS-873. At 60 min of incubation, progesterone or A23187 were added to stimulate acrosome reaction. The percentage of live sperm that undergo acrosome reaction is shown. Results are expressed as the mean \pm SEM of at least 6 independent experiments. * $p < 0.05$ represent statistical significance versus control (Pg). One-way ANOVA was performed in combination with Dunnett 's multiple comparisons test.

B) IVF was performed using sperm that were preincubated with DMSO (control) or 20 μ M NMS-873. Data represents the mean \pm SEM of at least 4 independent experiments. * $p < 0.05$ and ** $p < 0.01$ represents statistical significance vs conditions without progesterone. Paired t-test was performed.