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Cyclic nucleotide phosphodiesterase 3A–deficient mice as a model of female infertility

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Since cAMP blocks meiotic maturation of mammalian and amphibian oocytes in vitro and cyclic nucleotide phosphodiesterase 3A (PDE3A) is primarily responsible for oocyte cAMP hydrolysis, we generated PDE3A-deficient mice by homologous recombination. The Pde3a−/− females were viable and ovulated a normal number of oocytes but were completely infertile, because ovulated oocytes were arrested at the germinal vesicle stage, and therefore, could not be fertilized. Pde3a−/− oocytes lacked cAMP-specific PDE activity, contained increased cAMP levels, and failed to undergo spontaneous maturation in vitro (up to 48 hours). Meiotic maturation in Pde3a−/− oocytes was restored by inhibiting protein kinase A (PKA) with adenosine-3′,5′-cyclic monophosphorothioate, Rp-isomer (Rp-cAMPS) or by injection of protein kinase inhibitor peptide (PKI) or mRNA coding for phosphatase CDC25, which confirms that increased cAMP-PKA signaling is responsible for the meiotic blockade. Pde3a−/− oocytes that underwent germinal vesicle breakdown showed activation of MPF and MAPK, completed the first meiotic division extruding a polar body, and became competent for fertilization by spermatozoa. We believe that these findings provide the first genetic evidence indicating that resumption of meiosis in vivo and in vitro requires PDE3A activity. Pde3a−/− mice represent an in vivo model where meiotic maturation and ovulation are dissociated, which underscores inhibition of oocyte maturation as a potential strategy for contraception.

Introduction
Although competent to complete meiosis, mammalian oocytes are physiologically arrested in prophase I (prophase of the first meiotic division) until shortly before ovulation. Through protein kinase A-catalyzed (PKA-catalyzed) phosphorylation of unidentified proteins, cAMP prevents activation of maturation-promoting factor (MPF) and MAPK signaling in oocytes and inhibits the spontaneous maturation that occurs in vitro, thus maintaining meiotic arrest (1–5). It is unclear, however, whether the same mechanisms of meiotic arrest operate in vivo during each reproductive cycle, where preovulatory gonadotrophin triggers resumption of meiosis and progression through the second meiotic division until metaphase II. Only oocytes arrested in metaphase II can be fertilized.

Cyclic AMP apparently plays an important role in maintaining meiotic arrest in mammalian oocytes. The interplay of signals arising in both follicle cells and oocytes themselves (1, 2, 4, 6, 7) regulates synthesis and degradation of oocyte cAMP via adenylyl cyclases and cyclic nucleotide phosphodiesterases (PDEs), respectively. Cyclic AMP may enter oocytes from adjacent cumulus cells via gap junctions (2, 6). Recent studies in rodents, however, demonstrated that active oocyte adenylyl cyclase contributes to meiotic blockade (8), which can be released by microinjection of oocytes with antibodies that inactivate Gs, the heterotrimeric GTP-binding protein that activates adenylyl cyclase (9). These studies imply that generation of intra-oocyte cAMP is sufficient to maintain meiotic arrest. In Xenopus oocytes, meiotic arrest involves PKA-induced phosphorylation of protein phosphatase CDC25 (10), and in oocytes from sterile female Cdc25b−/− mice, which are arrested in prophase I, microinjection of CDC25B reinitiated meiosis (11), suggesting that phosphorylation/dephosphorylation of critical effectors is important in oocyte maturation (1, 3, 5, 10, 11).

PDEs belong to a complex and diverse superfamily of at least 11 structurally related, highly regulated, and functionally distinct gene families (PDE1–PDE11), which differ in their primary structures, affinities for cAMP and cGMP, responses to specific effectors, sensitivities to specific inhibitors, and regulatory mechanisms (12). Most PDE families comprise more than one gene, which generate multiple protein products via alternative mRNA splicing or utilization of different promotors and/or transcription initiation sites. The two PDE3 subfamilies, PDE3A and PDE3B, are encoded by closely related genes (13). PDE3A is relatively highly expressed in oocytes, platelets, and cardiac, vascular, and airway myocytes; PDE3B, in adipose tissue, liver, and pancreas, as well as cardiovascular tissues (13–16). PDE3 inhibitors increase myocardial contractility, inhibit platelet aggregation, and enhance vascular and airway smooth muscle relaxation (12, 13). Activation of PDE3B is thought to be important in the antilipolytic and antiglycogenolytic actions of insulin, as well as in IGF1- and leptin-induced inhibition of cAMP-stimulated secretion of insulin from pancreatic islets (16–19). Other studies suggest that, in pancreatic islets, PDE3B regulates intracellular cAMP pools that modulate glucose- and glucagon-like peptide-1–stimulated (GLP1-stimulated) insulin release (20, 21).

Oocyte cAMP hydrolysis is thought to be primarily accomplished by PDE3A (22, 23), and selective PDE3 (not PDE4 or PDE5) inhibi-
tors block meiotic resumption and oocyte maturation in *Xenopus* oocytes, culturedmurine, macaque, and human oocytes, and in vivo in cycling rats (in the latter, without affecting ovulation) (1, 22–27). Moreover, murine oocyte maturation, induced by microinjection of an active PDE (3), is associated with decreases in cAMP (1, 30, 31). We previously reported that heterologous expression of PDE3A in *Xenopus* oocytes induced activation of cAMP hydrolysis (via an enzyme with properties of PDE3) both signal by reducing adenyl cyclase and IGF1-induced activation of cAMP hydrolysis (32).

To understand specific physiological roles of PDE3A, and to elucidate the role of cyclic nucleotides in oocyte maturation and female infertility, we generated mice deficient in PDE3A. These PDE3A-deficient mice represent the first murine genetic model demonstrating that disruption of cAMP signaling leads to meiotic arrest of oocytes and female infertility, that the processes of meiotic maturation and ovulation are dissociated, and that PDE3A is essential for oocyte maturation and subsequent fertilization.

**Results**

To establish a model for delineating PDE3A functions, we generated PDE3A-deficient mice by homologous recombination in embryonic stem cells (Figure 1). Since multiple PDE3A isoforms, generated from the single *Pde3a* gene, are found in cardiovascular tissues (33, 34), *Pde3a*−/− mice were generated by a targeted disruption of exon 13, which encodes a portion of the PDE3A catalytic domain (Figure 1). As seen in Figure 1, mouse *Pde3a* (MPde3a) exon 13, encoding a portion of the second putative metal-binding site in the center of the PDE3A catalytic domain (Figure 1A), was replaced by the neomycin phosphotransferase II (NPTII) coding sequence (Figure 1B). Homologous recombination was confirmed by Southern blots (Figure 1C), or PCR of genomic DNA (Figure 1C, bottom). The mRNA of the rearranged gene, however, was transcribed and translated as shorter mutant PDE3A forms (Figure 1, D and E), lacking that portion of the catalytic domain encoded by exon 13 (Figure 1D). The immunoreactive, smaller, mutant PDE3A forms are expressed at low levels in *Pde3a*−/− mice and are much less abundant than intact WT PDE3A (Figure 1E). Consistent with PDE3A being the predominant PDE in murine oocytes (22, 23) and PDE3B being absent from oocytes (14, 15), and with the mutant form of PDE3A being catalytically inactive, PDE3 activity was undetectable in isolated *Pde3a*−/− oocytes (Figure 2A), and cAMP content was greater in denuded *Pde3a*−/− than in *Pde3a*−/+ oocytes (*P* = 0.003) (Figure 2B).

Because cells contain representatives of several PDE gene families, PDE3 activity in different tissue homogenates was measured (35) as that fraction of total PDE activity inhibited by 1 μM cilostamide, a specific PDE3 inhibitor. Cilostamide, however, inhibits both PDE3A and PDE3B isoforms with similar potency. As seen in Figure 2C, there was also almost complete loss of PDE3 activity in homogenates from *Pde3a*−/− lungs, indicating that PDE3A is the predominant PDE expressed in lung. PDE3 activity was also markedly reduced in homogenates from *Pde3a*−/− hearts. Consistent with relatively high expression of PDE3B in liver and adipose tissue, PDE3 activities were only slightly lower in those tissues from *Pde3a*−/− mice than from WT *Pde3a*−/+ mice (Figure 2C). Further investiga-
tion by gel filtration chromatography demonstrated that PDE3B could account for the residual low levels of PDE3 activity detected in lung homogenates prepared from Pde3a–/– mice (Figure 2D).

Mice lacking PDE3A were viable and exhibited no obvious growth or developmental deficiencies. Male Pde3a–/– mice were fertile and produced viable offspring, whereas female Pde3a–/– mice were completely sterile, whether mated to Pde3a–/– or Pde3a+/+ males (Table 1). Female Pde3a–/– mice were also fertile, which indicates that the residual, catalytically inactive, shorter mutant PDE3A forms (cf. Figure 1, D and E) did not interfere with oocyte function by acting as “dominant negative” PDE3A forms. Pde3a–/– ovarian structure and morphology were unremarkable, with a normal complement of corpora lutea and antral follicles, which suggests normal folliculogenesis and ovulation (Figure 3, A–H). As also seen in Figure 3, I–L, after injection of female mice with pregnant mare serum gonadotropin (PMSG) and human chorionic gonadotropin (hCG), cumulus expansion was normal and observed in ovarian follicles of both Pde3a+/+ and Pde3a–/– mice. Whereas the Pde3a+/+ oocyte progressed to metaphase, as evidenced by the presence of a metaphase spindle (Figure 3, I and J), the Pde3a–/– oocyte remained arrested in meiotic prophase I, as demonstrated by the persistence of the nuclear germinal vesicle (GV), classified as the GV stage (Figure 3, K and L). Although Pde3a–/– mice exhibit normal mating behavior and spontaneous and PMSG/hCG-stimulated ovulation (Table 2), nonfertilized Pde3a–/– oocytes retrieved from the ampulla (Table 2) also were arrested in meiotic prophase I (GV stage), as demonstrated by the persistence of the nuclear germinal vesicle (Figure 4B). After mating, two-cell embryos were collected from oviducts of Pde3a+/+ females (Table 2 and Figure 4A), whereas immature GV-stage oocytes with sperm in the perivitelline space were retrieved from Pde3a–/– oviducts (Figure 4B). Thus, sterility of Pde3a–/– females,
which apparently exhibit normal estrous cycles, ovulation, and mating behavior, is most likely related to the inability of Pde3a−/− oocytes to resume meiosis in response to the normal luteinizing hormone surge or exogenously administered gonadotropins.

As shown in Figure 4, C–H, immediately after harvesting from ovarian follicles (time 0), both Pde3a+/+ (Figure 4C) and Pde3a−/− (Figure 4D) oocytes contained intact GV, an interphase-like distribution of microtubules, and perinucleolar condensation of chromatin (surrounded nucleolus [SN] conformation). After removal from follicles, Pde3a+/+ oocytes, but not Pde3a−/− oocytes, resumed meiosis and completed germinal vesicle breakdown (GVBD) by approximately 3 hours, which was followed by extrusion of polar body (PB) (Figure 5, A–C). After 20 hours in culture, Pde3a−/− oocytes exhibited a well-defined spindle apparatus with condensed and aligned chromosomes typical of metaphase II (Figure 4E). In contrast, even after 20 hours, denuded Pde3a−/− oocytes did not resume meiosis in vitro and remained blocked at the GV stage, with an interphase pattern of dispersed tubulin and partially condensed chromatin (Figure 4F). MPF and MAPK were activated during spontaneous maturation (at 18 hours) of cultured Pde3a+/+ oocytes, but not Pde3a−/− oocytes (Figure 5D). Moreover, meiosis did not resume in cumulus-oocyte complexes cultured up to 12 hours with or without follicle-stimulating hormone (FSH), although the hormonal treatment produced normal expansion of cumulus cells (data not shown). Thus, complete blockade of spontaneous or gonadotropin-induced meiotic resumption occurs in Pde3a−/− oocytes.

Consistent with the absence of oocyte PDE3A, cAMP levels were higher in Pde3a−/− than in Pde3a+/+ oocytes (cf. Figure 2B). Given the proposed role of cAMP and PKA in maintenance of meiotic arrest (1–5), in vitro treatment of Pde3a+/+ oocytes with PKA inhibitors Rp-8-Br-cAMPS or adenosine-3′,5′-cyclic monophosphorothiate, Rp-isomer (Rp-cAMPS) induced resumption of meiosis (Figure 5E).

As expected, incubation of Pde3a+/+ oocytes with cilostamide, a PDE3 inhibitor, blocked GVBD (Figure 5F). Treated Pde3a−/− oocytes exhibited activated MPF and MAPK (Figure 6A) and were competent for fertilization in vitro (Figure 6B). Microinjection of PKI or protein phosphatase cdc25b mRNA also initiated resumption of meiosis in Pde3a−/− oocytes (Figure 6C).

Discussion

The PDE gene superfamily apparently encodes more than 50 PDE proteins (12). Some cells are relatively enriched in specific PDEs; these include oocytes (22, 23) and retinal rods and cones (12), in which PDE3A and PDE6, respectively, are the predominant PDEs. Most cells, however, contain representatives of at least several PDE gene families and different subfamily isoforms from the same gene family, but in different amounts, proportions, and subcellular locations (12). Family-specific PDE inhibitors can pharmacologically define functions of individual PDEs and, in cells containing multiple PDEs, dissect out effects of specific PDEs in regulating specific signaling pathways and cellular functions. For example, PDE3 inhibitors, but not PDE4 inhibitors, increase insulin secretion in cultured pancreatic cells ( HIT-T15 or HIT cells), despite greater effects of PDE4 inhibitors on cAMP content in these cells (17, 18). In the female reproductive system, differential cellular expression of PDE4 and PDE3 isoforms in granulosa cells and oocytes, respect-

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**Table 1**

<table>
<thead>
<tr>
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<th>M</th>
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<th>Average litter sizea</th>
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<td>7.9 ± 2.9</td>
<td>10</td>
</tr>
<tr>
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<td>WT</td>
<td>he</td>
<td>6.1 ± 2.0</td>
<td>7</td>
</tr>
<tr>
<td>KO</td>
<td>he</td>
<td>KO</td>
<td>7.2 ± 2.0</td>
<td>27</td>
</tr>
<tr>
<td>he</td>
<td>he</td>
<td>KO</td>
<td>0 ± 0</td>
<td>4</td>
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</table>

^a mean ± SEM, n = 4–27 litters, resulting from matings of C57BL/6J-129/SvJ Pde3a+/+ (WT) and Pde3a−/− (KO) mice; ^p indicates numbers of litters. Identical results, i.e. infertility of KO females, were observed with Pde3a−/− and Pde3a+/+ mice resulting from crossing Black Swiss and C57BL/6J-129/SvJ mice. he, heterozygous.

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**Figure 3**

Histology of ovaries from Pde3a+/+ and Pde3a−/− mice. (A–H) H&E sections of ovaries harvested from C57BL/6J-129/SvJ females (3 months old), Pde3a+/+ (A and C) or Pde3a−/− (B and D), or from females (5 weeks old) after induction of superovulation as described in Methods, Pde3a+/+ (E and G) or Pde3a−/− (F and H). Magnification, A, B, E, and F: ×4; C, D, G, and H: ×20. Pde3a+/+ females exhibited normal ovaries and follicles, either in normal condition or following superovulation. (I–L) Pde3a+/+ or Pde3a−/− females were injected with PMSG and hCG as described in Methods; 9 hours after injection with hCG, H&E sections were prepared. Note the expanded cumulus, which has lost contact with the follicle wall in both genotypes. Whereas the Pde3a−/− oocytes are in metaphase (with an evident spindle) (I and J), the immature Pde3a−/− oocytes exhibit an intact nuclear membrane (GV stage) (K and L).
tively, seems to be important in the integrating signals involved in luteinizing hormone (LH) induction of oocyte maturation, which apparently involves opposing effects of cAMP in somatic and germ cells (1, 15). Actions of PDE4 inhibitors in follicle cells enhance effects of LH on oocyte maturation, whereas PDE3 inhibitors act directly in oocytes, increasing cAMP and suppressing meiotic progression (1, 15). In PDE4D KO mice, however, the rate of ovulation and, consequently, fertility, is reduced (36). Other reports further support the view that multiplicity of PDEs does not serve merely a survival or protective function, but that differentially regulated and targeted PDEs are critical for selective compartmentalization and modulation of cyclic nucleotide signals and responses (37, 38).

Table 2

Absence of fertilization in Pde3a<sup>−/−</sup> oocytes following natural or induced ovulation

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Ovulated eggs</th>
<th>Embryos</th>
<th>Oocytes</th>
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<tr>
<td>WT</td>
<td>Natural ovulation (C57BL/6J-129/SvJ)</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>KO</td>
<td>28</td>
<td></td>
<td></td>
</tr>
<tr>
<td>WT</td>
<td>Induced ovulation (C57BL/6J-129/SvJ)</td>
<td>90</td>
<td>90</td>
</tr>
<tr>
<td>KO</td>
<td>122</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<sup>a</sup>n indicates numbers of female mice. Natural ovulation: Females (C57BL/6J-129/SvJ) of the indicated genotypes (n = 1 or 4) were individually caged with WT males and sacrificed the morning after mating (identified by the presence of vaginal plugs). Ovulated cells, i.e., embryos or oocytes, were harvested from the oviducts of either Pde3a<sup>−/+</sup> or Pde3a<sup>−/−</sup> mice, pooled, and counted. Induced ovulation: Three females (C57BL/6J-129/SvJ) of the indicated genotypes were induced to ovulate by a priming injection of PMSG followed 46 hours later by an injection of hCG, as described in Methods. On the day of hCG injection, the females were individually caged with WT males and sacrificed the morning after mating. Embryos or oocytes were harvested from the oviducts of either Pde3a<sup>−/+</sup> or Pde3a<sup>−/−</sup> mice, pooled, and counted. The same results were obtained when this experiment was repeated with Pde3a<sup>−/+</sup> and Pde3a<sup>−/−</sup> mice resulting from crossing Black Swiss and C57BL/6J-129/SvJ mice.

Figure 4

Maturation of Pde3a<sup>−/+</sup> and Pde3a<sup>−/−</sup> oocytes in vivo and in vitro. (A and B) Nomarski images of Pde3a<sup>−/+</sup> embryos and nonfertilized, ovulated Pde3a<sup>−/−</sup> oocytes. Following induced ovulation, and 24 hours after mating, Pde3a<sup>−/+</sup> embryos and nonfertilized, ovulated Pde3a<sup>−/−</sup> oocytes were harvested from oviducts of females as described in Methods. (A) Two-cell–stage embryos were observed in oviducts from Pde3a<sup>−/+</sup> females. (B) Immature Pde3a<sup>−/−</sup> oocytes were not fertilized and exhibited clear intact nuclear membranes (GV) (thin arrows) and nucleoli (n). Sperm (arrowheads) traversed the zona pellucida (zp) but did not fuse with the plasma membranes of immature GV stage oocytes. (C–H) Nomarski images and immunofluorescent staining of oocytes from Pde3a<sup>−/+</sup> and Pde3a<sup>−/−</sup> mice. Pde3a<sup>−/+</sup> (C) and Pde3a<sup>−/−</sup> (D) oocytes were fixed immediately after isolation. Pde3a<sup>−/+</sup> (E and G) and Pde3a<sup>−/−</sup> (F and H) oocytes were incubated for 20 hours in maturation medium without (E and F), or with (G and H), 5 mM Rp-8-Br-cAMPS. In each row from left to right are shown Nomarski images, antitubulin immunofluorescent staining (green), DAPI immunofluorescent staining (blue), and merged images of anti-tubulin and DAPI staining. In these oocytes the chromatin nucleolar rim around the nucleoli (red arrows) and the interphase-like microtubules can be seen. After 20 hours, metaphase II–blocked oocytes show the chromosomes aligned at the metaphase plate (yellow arrows), microtubule spindle formation (green arrows), and PBs (orange arrow). For technical reasons, in F, two different Pde3a<sup>−/−</sup> oocytes were used for Nomarski imaging and immunofluorescence staining.
plays a unique role, which is apparently not replaced or functionally compensated for by other PDE gene families or by the other PDE3 family member, PDE3B. Inactivation of the \textit{Pde3a} gene results in a phenotype characterized by female infertility. Despite normal ovarian structure, folliculogenesis, ovulation, and mating behavior, female PDE3A KO mice are sterile, most likely due to specific and direct effects of the absence of PDE3A in oocytes — i.e., inhibitory effects of elevated cAMP on meiotic progression and oocyte maturation and, consequently, the competency of oocytes for fertilization. Additional studies with PDE3A KO mice will undoubtedly prove useful in elucidating other physiological roles of PDE3A.

Targeted disruption of Pde4 genes has also similarly generated mice with unique phenotypes (36, 39–41). Analyses of these phenotypes support the concept that different PDEs are functionally nonredundant and regulate different cAMP/cGMP pools and responses. Although increases in cAMP and activation of PKA are crucial in the induction and maintenance of meiotic arrest, the precise targets and downstream effectors of cAMP signalling have not been completely defined (1, 42). Recent studies suggest that PKA may affect two parallel pathways that regulate activation of MPF (3, 10, 42, 43). PKA apparently blocks translation of Mos (the oocyte MAPK), perhaps by inhibiting polyadenylation of its mRNA (43). In frog oocytes, Mos activates MAPK, leading to inhibition of MYT1, a kinase that phosphorylates and inhibits MPF (3, 10, 42, 43). In addition, PKA-catalyzed phosphorylation of CDC25 may result in its sequestration by 14-3-3 and thus negatively regulate CDC25-induced activation of MPF (10).

The rapid reversal of cAMP/PKA-induced meiotic arrest by PKA inhibitors in Pde3a−/− oocytes (cf. Figures 5 and 6) is consistent with the reported activation of PDE3A (23) and decrease in cAMP (28, 29) that apparently precede spontaneous or hCG-induced maturation of murine oocytes. Thus, a brief period of reduced PKA activity (via PKA inhibition in Pde3a−/− oocytes or activation of PDE3A and reduction in cAMP and PKA activity in murine

**Figure 5**

Regulation of maturation of cultured Pde3a+/+ and Pde3a−/− oocytes. (A–C) Time course of spontaneous maturation. (A) Denuded Pde3a+/+, not Pde3a−/−, oocytes undergo spontaneous maturation (demonstrated by GVBD) within 3 hours in vitro. (B and C) After 20 hours, Pde3a+/+ (B), but not Pde3a−/− (C), oocytes exhibit GVBD and PBs. (D) Absence of MAPK and MPF activation in Pde3a−/− oocytes. MAPK and MPF, assayed at indicated times as described in Methods, were activated (18 h) in Pde3a+/+, but not Pde3a−/−, oocytes. (E and F) Effects of PKA inhibitors on maturation of Pde3a+/+ and Pde3a−/− oocytes. (E) Pde3a+/+ (with 10 µM cilostamide) and Pde3a−/− oocytes were incubated for 24 hours without or with 5 mM Rp-cAMPS before evaluation of GVBD. In the absence of Rp-cAMPS, spontaneous maturation was blocked in both Pde3a+/+ and cilostamide-treated Pde3a−/− oocytes. GVBD and meiosis were reinitiated to the same extent by Rp-cAMPS in Pde3a+/+ and cilostamide-treated Pde3a−/− oocytes. (F) Pde3a+/+ and Pde3a−/− oocytes were incubated for the indicated times with 5 mM Rp-8-Br-cAMPS, washed twice in a drop (50 µl) of maturation medium, and incubated without Rp-8-Br-cAMPS. After 24 hours, all oocytes were assessed for GVBD and PB formation. “Total oocytes” indicates the numbers incubated under each condition. Within 1 hour of treatment of Pde3a−/− oocytes with Rp-8-Br-cAMPS, meiosis was reinitiated, as monitored by the percent decrease in GV, and increase in GVBD and PB. Data at 0, 1, 3, and 24 hours represent mean percent ± one half of the range, in two separate experiments.
Taken together, these experiments strongly suggest that absence of PDE3A in oocytes leads to high levels of cAMP, persistent activation of PKA, and phosphorylation of critical regulatory effectors, resulting in inhibition of MPF/MAPK signals involved in resumption of meiosis and subsequent development of competency for fertilization (Figure 7). Under these conditions, the physiological signal that triggers meiosis in vivo is ineffective, which underscores the critical role of PDE3A in meiotic resumption and female fertility. Thus, Pde3a–/– mice provide a novel genetic model for studies of female sterility and present the possibility that some idiopathic infertility in women may be the result of mutations that cause inactivation of this gene. More importantly, the apparently reversible, perhaps "physiological," nature of the meiotic arrest in Pde3a–/– oocytes clearly suggests — along with earlier reported in vivo and in vitro effects of PDE3 inhibitors on meiotic progression and oocyte maturation (26, 27) — that PDE3A could be an excellent potential target for contraceptives (Figure 7). It should be pointed out that ovulation and oocyte maturation are completely dissociated in the Pde3a–/– mouse, which emphasizes the concept that oocyte maturation can be targeted independently of somatic cells and without affecting the hormonal milieu. At the present time, however, use of PDE3 inhibitors, such as cilostamide or clostazol, as contraceptives is not feasible, since inhibitors specific for PDE3A or PDE3B isoforms have not been developed, and methods to selectively target oocyte PDE3A are not available. Nevertheless, as outlined in Figure 7, Pde3a–/– mice provide a valuable model in which to dissect and reconstitute signaling pathways downstream of cAMP that regulate meiosis in oocytes (and perhaps division and proliferation in other cAMP-responsive cells). It is also conceivable that some of these downstream effectors and pathways will offer more accessible and efficacious contraceptive targets.

Methods

Generation of Pde3a–/– mice. Pde3a–/– mice were generated by a targeted disruption of exon 13, which encodes a portion of the second putative metal-binding site in the center of the PDE3A catalytic domain. The pKO Scrambler NTK Vector-1903 (Stratagene, La Jolla, California, USA) was used to construct the replacement vector. An approximately 16-kb SalI genomic DNA fragment, containing five putative exons (exons 10–14) that encode much of the catalytic domain of PDE3A was ligated into the SalI site of the vector. This vector was used to generate a C57Bl/6 ES clone by standard techniques. Southern analysis was done on DNA prepared from ES clones and positive clones were used for injection. Male and female PDE3A–/– mice provided fertile offspring and do not exhibit overt abnormalities, indicating that disruption of PDE3A is not lethal. The protein levels of PDE3A were assessed in a variety of tissues from PDE3A–/– mice using Western blot analysis. A-kinase–anchoring proteins (AKAPs) (44, 45) might compartmentalize, and thereby regulate actions of, PKA and other effector molecules, i.e., protein kinases and phosphatases and other scaffolding proteins. Whether PDE3A is physically targeted to these putative spatially segregated signaling modules (similar to the association of PDE4 isoforms with AKAP in regulating ryanoamine-sensitive channels [ref. 46] or with β-arrestin in downregulation of the β-adrenergic receptor [ref. 47]), and/or regulates their signaling output, is uncertain. These considerations are also consistent with recent studies indicating that some of the inhibitory effects of PKA on meiotic arrest may be independent of kinase activity and perhaps mediated via its direct interaction with and sequestration of critical regulatory effectors rather than via phosphorylation of these molecules (48).
The resulting chimera was mated with C57BL/6J to produce the F₁ generation blastocysts and implanted into pseudopregnant females. Those positive ES cell clones, identified by Southern blots and PCR, were microinjected into C57BL/6J-129/SvJ mice, respective to the orientation of the Neo sequence introduced a selectable marker, and replaced by the NPTII coding sequence. Positive ES cell clones, called fragment A) and fragment B, with containing the NPTII coding sequence (3.5 kb, called fragment B) were sequentially cloned into the two polylinker sites of the pKO vector. This vector was linearized and electroporated into C57BL/6J blastocysts and implanted into pseudopregnant females. Our results with PKA inhibitors, such as Rp-8-Br-cAMPS and PKI, are consistent with PKA playing a major role in maintaining meiotic arrest at prophase I and thereby preventing meiotic progression to metaphase II and consequent development of competency for fertilization. The apparently reversible nature of the meiotic arrest in Pde3a−/− oocytes clearly suggests, along with reported effects of PDE3 inhibitors, such as cilostamide, on oocyte maturation (22–27), that oocyte PDE3A could represent an excellent potential target for contraceptives.

**Figure 7**

Role of PDE3A in oocyte maturation and female infertility. Pde3a+/− mice represent what we believe to be the first genetic model indicating that PDE3A is necessary for meiotic resumption in mouse oocytes and that, in the absence of oocyte PDE3A, increased cAMP and the consequent disruption of cAMP-signaling can lead to female infertility, presumably via PKA-catalyzed phosphorylation of key regulatory effectors that result in inhibition of MPF/MAPK-induced oocyte maturation. The vector was linearized and electroporated into C57BL/6J blastocysts and implanted into pseudopregnant females. Our results with PKA inhibitors, such as Rp-8-Br-cAMPS and PKI, are consistent with PKA playing a major role in maintaining meiotic arrest at prophase I and thereby preventing meiotic progression to metaphase II and consequent development of competency for fertilization. The apparently reversible nature of the meiotic arrest in Pde3a−/− oocytes clearly suggests, along with reported effects of PDE3 inhibitors, such as cilostamide, on oocyte maturation (22–27), that oocyte PDE3A could represent an excellent potential target for contraceptives.

of MPDE3A, was isolated from a 129/SvJ FIXII mouse genomic library (Stratagene), using a [32P]-labeled probe generated from a cDNA clone containing the catalytic domain of rat PDE3A. A 7.0-kb ScaI DNA fragment containing putative exons 11–14 was subcloned from this original clone into pUC18 (ScaI site). This 7.0-kb genomic subclone includes an approximately 1.5-kb Acl restriction fragment that contains exon 13. Two segments, ScaI-Aacl (2.0 kb, called fragment A) and Aocl-ScaI (3.5 kb, called fragment B) were amplified by PCR and purified. Fragment A was digested with EcoRI and EcoRI fragments AocI-SacI (3.5 kb, called fragment B) were amplified by PCR and purified. Fragment A was digested with EcoRI and EcoRI fragments AocI-SacI (3.5 kb, called fragment B) were amplified by PCR and purified. Fragment A was digested with EcoRI and EcoRI fragments AocI-SacI (3.5 kb, called fragment B) were amplified by PCR and purified. Fragment A was digested with EcoRI and EcoRI fragments AocI-SacI (3.5 kb, called fragment B) were amplified by PCR and purified. Fragment A was digested with EcoRI and EcoRI fragments AocI-SacI (3.5 kb, called fragment B) were amplified by PCR and purified. Fragment A was digested with EcoRI and EcoRI fragments AocI-SacI (3.5 kb, called fragment B) were amplified by PCR and purified. Fragment A was digested with EcoRI and EcoRI fragments AocI-SacI (3.5 kb, called fragment B) were amplified by PCR and purified.
GVBD stage; and metaphase II–arrested oocytes, with a PB, were classified as PB stage (27). After spontaneous or induced ovulation, the ovulated cells were examined; those at the two-cell stage were classified as embryos, whereas those still exhibiting a clear GV were defined as GV stage ovulated oocytes. Progression of meiotic maturation of denuded oocytes in vitro was scored by monitoring GVBD with an inverted microscope fitted with a Hoffman contrast lens.

**Immunofluorescence.** Twenty to 30 oocytes were fixed for 1 hour at room temperature (rt) in 2% paraformaldehyde in PBS, washed 5 times in blocking solution (PBS, pH7.4, containing 0.3% BSA [Sigma-Aldrich] and 100 mM glycine [ICN Biomedicals Inc., Costa Mesa, California, USA]), permeabilized with blocking solution containing 0.1% Triton X-100 (Sigma-Aldrich), washed 3 times in blocking solution, and incubated 1 hour (rt) under embryo-tested mineral oil (Sigma-Aldrich) in 0.05 ml of anti-α-tubulin antibody (Accurate Chemical and Scientific Corp., Westbury, New York, USA), which had been diluted 1:100 in blocking solution. After five washes in blocking solution, the oocytes were incubated for 1 hour (rt) in 0.05 ml of the secondary, Alexa Fluor 488–conjugated anti-rat antibody (Molecular Probes, Eugene, Oregon, USA), and washed 5 times. After 10 minutes in PBS, pH 8.1/glycerol (1:1 vol/vol), the oocytes were mounted in mounting medium (DAPI; Vectashield, Burlingame, California, USA) using a Hybaid Easyscale frame (Continental Lab Products, San Diego, California, USA) with regular glass slides and coverslips and viewed with a confocal microscope (Leica Microsystems, Heidelberg, Germany).

**PDE activity measurements.** Denuded oocytes from Pde3a+/− and Pde3a−/− mice (20 for each PDE assay) were collected within 30 minutes after follicle puncture in Leibowitz’s L-15 medium (Invitrogen Corp., Grand Island, New York, USA) with 5% FBS and 100 IU/ml penicillin/streptomycin, washed, and frozen in PBS (3 µl). Immediately before assay, 20 µl of IPA buffer (10 mM sodium phosphate buffer, pH 7.2, 50 mM NaF, 150 mM NaCl, 2 mM EDTA, 5 mM mercaptoethanol, 30 mM pyrophosphate, 3 mM benzamidine, 5 µg/ml leupeptin, 20 µg/ml pepstatin, 2 mM PMSF, 1 µM microcystin, and 0.5% Triton X-100) were added. Samples were centrifuged at 12,000 g at 4°C for 30 minutes, supernatants collected, and PDE activity was measured for 20 minutes.

After sacrifice of mice by asphyxiation (CO2), heart, lung, liver, and fat pads were dissected from 3–3.5-month-old C57BL/6J-129/SvJ males, quickly washed in ice-cold PBS, and homogenized (5 ml/g tissue) in buffer (50 mM HEPES, pH 7.5 [Sigma-Aldrich], 25 mM KCl [Sigma-Aldrich], 5 mM MgCl2 [Quality Biological Inc., Gaithersburg, Maryland, USA], protease inhibitor cocktail tablets [Roche, Nutley, New Jersey, USA]) with a rotor-stator homogenizer (Omni International, Marietta, Georgia, USA) at 30,000 rpm (15 to 90 seconds, on ice), followed by homogenization (on ice, 20 strokes in a glass Dounce tissue grinder [B pestle; Thomas Scientific, Swedesboro, New Jersey, USA]) and sonification (on ice, 20 pulses, 40% duty cycle, output scale 4). NP-40 (Calbiochem) (1% final) was then added and samples incubated (on ice, 1 hour), before centrifugation (15,000 g, 20 minutes, 4°C). Supernatants (total tissue lysates) were used for PDE assays (35) or gel filtration chromatography (FPLC-superose 12; AKTA FPLC System; Amersham-Pharmac Ta Biotech Inc., Piscataway, New Jersey, USA). For PDE assays, samples (usually 5–15 µg protein in 100 µl) were incubated for 10 minutes in a total volume of 0.3 ml containing 50 mM HEPES, pH 7.4, 100 µM EGTA, 8.3 mM MgCl2, and [3H]-cAMP. PDE3 activity is that portion of total PDE activity inhibited by 1 mM cistolamide (Calbiochem), a specific PDE3 inhibitor. Inhibitor vehicle, DMSO (Sigma-Aldrich), added in equal quantities to samples without inhibitors, did not alter PDE activity. Protein was determined by BCA protein assay (Pierce, Rockford, Illinois, USA).

**Western blots.** Samples of solubilized tissue homogenates (~100 mg protein) were subjected to SDS-PAGE (4–12% Tris-Glycine gels [Invitrogen Corp., Carlsbad, California, USA]). After electrophoretic transfer, nitrocellulose membranes were blocked overnight at 4°C in T-PBS (0.05% Tween 20 [Bio-Rad Laboratories, Hercules, California, USA] in PBS) containing 5% non-fat dry milk (Bio-Rad Laboratories) and 1% BSA, incubated 1–4 hours with the indicated primary antibody diluted in T-PBS/5% milk, washed and incubated 1 hour with HP-conjugated secondary antibody against rabbit IgG (Amersham-Pharmacia Biotech Inc.) diluted 1:4000 in T-PBS. The signal was detected using ECL-Plus Detection System (Amersham-Pharmacia Biotech Inc.).

Primary antibodies, which were purified by chromatography on DEAE columns, included: rabbit IgG raised against a peptide corresponding to AAs 424–460 (regulatory domain) of human PDE3A, diluted 1:400 in T-PBS/5% milk; rabbit IgG raised against a peptide corresponding to AAs 2–18 (N-terminal domain) of rat PDE3B, diluted 1:200 in T-PBS/5% milk.

**Measurement of cAMP content.** Groups of 50 denuded oocytes were collected in L-15 medium, quickly washed, and frozen in PBS (3 µl). After thawing and extraction with 100 µl of 0.1% trichloroacetic acid in 95% ethyl alcohol, followed by centrifugation (800 g, 30 min, 4°C), the supernatant was dried, and the residue dissolved in PBS (100 µl). After acetylation, cAMP content was measured by radioimmunoassay (8).

**Maturation-promoting factor (MPF) and MAPK activity measurements.** Harvested oocytes were incubated for the indicated times prior to in vitro assay of MPF and MAPK as described (49), using histone H1 and myelin basic protein (MBP) as substrates, respectively. Briefly, 3 denuded oocytes (in 1.5 µl of maturation medium) were immediately frozen on dry ice in 3.5 µl of lysis buffer (7 µg/ml aprotinin, 7 µg/ml leupeptin, 7 mM p-nitrophenyl phosphate, 14 mM β-glycerophosphate, 70 µM sodium orthovanadate, and 3.5 mM EGTA) and stored at −80°C prior to assay. 10 µl final vol, 30 min at 30°C, 2 ng/ml histone [type III-S], 0.5 mg/ml MBP, and 500 µCi/ml [y-32-P]ATP [3,000 Ci/mmol; Amersham-Pharmacia Biotech Inc.]. Reactions were terminated and proteins separated by SDS-PAGE (15% gels), followed by gel drying and exposure to x-ray film.

**In vitro fertilization.** Denuded Pde3a+/− oocytes were exposed to 5 mM Rp-cAMPS in maturation medium for 2 hours, washed, and incubated overnight at 37°C in L-15 medium supplemented with 5% fetal bovine serum and 100 IU/ml penicillin/streptomycin. After 20 hours, oocytes at MII stage were incubated with capacitated sperm in modified KSOM embryo culture medium containing 5.6 mM glucose and 4 mg/ml BSA. After 4 hours, oocytes and attached sperm were transferred to fresh medium and further incubated at 37°C in 5% CO2 for 20 hours.

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