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The HDL receptor scavenger receptor, class B type I (SR-BI) controls the structure and fate of plasma HDL. Female SR-BI KO mice are infertile, apparently because of their abnormal cholesterol-enriched HDL particles. We examined the growth and meiotic progression of SR-BI KO oocytes and found that they underwent normal germinal vesicle breakdown; however, SR-BI KO eggs, which had accumulated excess cholesterol in vivo, spontaneously activated, and they escaped metaphase II (MII) arrest and progressed to pronuclear, MII, and anaphase/telephase III stages. Eggs from fertile WT mice were activated when loaded in vitro with excess cholesterol by a cholesterol/methyl-β-cyclodextrin complex, phenocopying SR-BI KO oocytes. In vitro cholesterol loading of eggs induced reduction in maturation promoting factor and MAPK activities, elevation of intracellular calcium, extrusion of a second polar body, and progression to meiotic stages beyond MII. These results suggest that the infertility of SR-BI KO females is caused, at least in part, by excess cholesterol in eggs inducing premature activation and that cholesterol can activate WT mouse eggs to escape from MII arrest. Analysis of SR-BI KO female infertility raises the possibility that abnormalities in cholesterol metabolism might underlie some cases of human female infertility of unknown etiology.


Significance

Production of functional sperm and eggs requires a complex process called meiosis. Meiosis in mouse and human eggs pauses at a stage called metaphase II (MII) arrest until fertilization by sperm. After fertilization, eggs released from MII arrest complete meiosis and develop into new individuals. In analyzing the female infertility of genetically altered mice, we discovered that excess cholesterol can trick mouse eggs into behaving as though they were fertilized (released from arrest), thus disrupting the normal synchrony between fertilization and completion of meiosis and rendering them dysfunctional. These findings suggest that abnormal cholesterol metabolism may contribute to some forms of human female infertility.


The authors declare no conflict of interest.

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Results

Our previous studies involving restoration of fertility by ovary transplantation and drug treatment suggested that the infertility of SR-BI KO (SR-BI<sup>−/−</sup>) female mice is unlikely to be caused by irreversibly impaired early oogenesis or abnormal folliculogenesis (2, 6, 7, 14) (Discussion). Rather, the infertility seems likely to be related to defects in oocyte growth, maturation, or activation. SR-BI KO females ovulate almost normal numbers of eggs (6). However, on their recovery from the oviduct, a substantial fraction of the ovulated eggs displays an abnormal, nonrefractile morphology characteristic of dead eggs (the criterion used to score for death here). On average, 4% of ovulated eggs from control SR-BI KO females accumulated excess cholesterol. Strikingly, we could recapitulate the apparently spontaneous activation of ovulated SR-BI KO females in eggs ovulated from fertile control mice (WT or SR-BI<sup>+/−</sup>) by treating those eggs with a cholesterol/methyl-β-cyclodextrin (MβCD) complex to load them with excess cholesterol (26). In addition, cholesterol loading induced a single spike in [Ca<sup>2+</sup>], reminiscent of that seen in ethanol-activated eggs (24, 25). These results establish that excess cholesterol loading provides a new approach to activating mammalian MII-arrested eggs and suggest that the infertility of SR-BI KO females is caused, at least in part, by excess cholesterol acting directly on the egg to induce premature activation. Analysis of SR-BI KO female infertility also raises the possibility that abnormalities in cholesterol metabolism might underlie some forms of human female infertility of unknown etiology.

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Here, we found that, within follicles of hormone-treated SR-BI KO females, oocytes apparently resumed meiosis normally. However, these cells spontaneously escaped MII arrest, exhibited reduced MPF and MAPK activities, and aberrantly progressed to haploid pronuclear, metaphase III, and anaphase/telophase III stages, similar to normal ovulated eggs after chemical (e.g., SrCl<sub>2</sub>) (22, 25) activation (Fig. 1). Compared with control eggs, the eggs of SR-BI KO females accumulated excess cholesterol. Strikingly, we could recapitulate the apparently spontaneous activation of ovulated SR-BI KO females in eggs ovulated from fertile control mice (WT or SR-BI<sup>+/−</sup>) by treating those eggs with a cholesterol/methyl-β-cyclodextrin (MβCD) complex to load them with excess cholesterol (26). In addition, cholesterol loading induced a single spike in [Ca<sup>2+</sup>], reminiscent of that seen in ethanol-activated eggs (24, 25). These results establish that excess cholesterol loading provides a new approach to activating mammalian MII-arrested eggs and suggest that the infertility of SR-BI KO females is caused, at least in part, by excess cholesterol acting directly on the egg to induce premature activation. Analysis of SR-BI KO female infertility also raises the possibility that abnormalities in cholesterol metabolism might underlie some forms of human female infertility of unknown etiology.

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a 50:50 C57BL/6J:129 genetic background (Materials and Methods). In some experiments, we used eggs from females on a WT, pure C57BL/6 background.

**Oocyte Follicular Growth.** Oocytes of the primordial follicle pool arrested in prophase I of meiosis (Fig. 1) are recruited to grow and differentiate throughout the mouse’s reproductive lifetime. As oocytes grow, their volume increases proportionally to the volume of the encasing follicle. In rodents, the relative oocyte to follicle diameter from intact follicles of unprimed females has been used as an indicator of follicular oocyte growth (27, 28). We mechanically isolated murine follicles from both SR-BI KO mice and fertile SR-BI*+/− controls at different stages of development from unprimed sexually immature (prepubertal) 3 wk-old and mature 6 wk-old females. The oocyte and follicle diameters were measured as described in Materials and Methods. Fig. 2 A and B shows that, at both ages, there were linear relationships between the oocyte and follicle diameters and that the slopes of these lines were virtually identical for the follicles and oocytes from SR-BI KO females and SR-BI*+/− controls. Thus, SR-BI deficiency does not seem to alter the growth of oocytes and their surrounding follicles.

**Meiotic Resumption and Extrusion of the First PB.** An important indicator of oocyte quality is the ability to resume and complete meiosis, a process called maturation (29). In mice, oocytes initiate meiosis during fetal development and progress through prophase I until the diplotene stage, where they enter primary arrest (Fig. 1). These primordial oocytes acquire the competence to resume meiosis during the late stages of growth and differentiation. In full-grown oocytes, primary arrest is maintained by the follicular soma and relieved before ovulation. Oocytes that resume meiosis undergo germinal vesicle breakdown (GVBD), assemble chromosomes on the first metaphase plate, and complete the first meiotic division, which is accompanied by extrusion of the first PB (Fig. 1). Oocytes then assemble the remaining single set of homologous chromosomes on the second metaphase plate and enter secondary arrest (MII arrest) (Fig. 1).

Full-grown, prophase I-arrested oocytes in the ovary can be triggered to resume meiosis and progress to MII and secondary arrest by removing them from their follicles. Thus, to investigate whether full-grown oocytes from the SR-BI KO mice and SR-BI*+/− controls were competent to resume meiosis, we isolated them from ovaries of unprimed females by follicular puncture and cultured them for 14 h. Resumption and completion of MI were scored by determining by microscopic examination GVBD and first PB extrusion, respectively (Materials and Methods). Table S1 shows that the fractions of SR-BI*+/− and SR-BI KO oocytes undergoing GVBD were similar (87% and 74%, respectively), and the fractions of oocytes extruding the first PB were also similar (78% and 64%, respectively). Fig. 2C shows that the rate of GVBD in oocytes from SR-BI KO animals was similar to that for SR-BI*+/− controls. These results indicate that the rate and extent to which SR-BI KO oocytes resume (GVBD) and complete MI (first PB extrusion) are similar to those of control, fertile oocytes, indicating that they are competent to resume meiosis.

**Spontaneous Progression Beyond MII Arrest.** After completion of MI, WT eggs, which still reside in the ovarian follicles, arrest at MII, at which point they are ovulated. The ovulated eggs in the oviduct remain arrested at MII until they are activated by sperm during fertilization, which triggers the completion of meiosis II and the extrusion of the second PB (Fig. 1). Eggs can also activate spontaneously (e.g., during aging) or artificially (e.g., by treatment with chemicals, such as SrCl2/EGTA, ionophores, or ethanol) (30–33). To study postovulatory events in eggs from SR-BI KO mice, we harvested morphologically live eggs from the oviducts of hormone-primed SR-BI KO and control SR-BI*+/− females either ~16 or ~20 h after hCG administration. We determined the meiotic stage of both control SR-BI*+/− and SR-BI KO eggs by examining their chromosome/nuclear and spindle organizations. Fig. 3 shows eggs that were stained using a whole-mount preparation technique and imaged by deconvolution microscopy to visualize the DNA (DAPI, blue), microtubules (anti-tubulin, green), nuclear envelope (anti-lamin B1, see Fig. S1), and centromeres (anti-centromere antibody, red). We present the 3D organization of chromosomes in the egg and its PB as a z stack of deconvolved images. After image collection and processing, we scored the meiotic state of each egg by determining the number of chromatids by counting centromeres (one chromatid per centromere) in the egg and its PB as well as scoring the morphology of the spindle and the presence or absence of the nuclear lamina. The results are presented in Table 1, and representative images are shown in Fig. 3.

The two leftmost columns in Fig. 3 show control SR-BI*+/− and SR-BI KO eggs at metaphase II contained 40 sister chromatids (20 pairs) aligned on a spindle (Fig. 3, two leftmost columns, spindles perpendicular to the plane of the image). The other SR-BI KO eggs were classified, based on their chromosomal and spindle morphologies, into five categories, for which in Fig. 3 schematic representations (top row) and representative images at several magnifications (rightmost five columns) are shown. In each case, progression beyond MII was evident from the segregation of sister chromatids (40 at MII), leaving 20 single chromatids.
most of the control SR-BI+/+ eggs (n = 8) were either arrested at MII (Fig. 3, first column) or at an earlier stage at ~16 h (93% of eggs scored) or ~20 h (89%) after hCG administration (induction of superovulation) (Table 1). In this experiment, a small fraction of the control SR-BI+/+ eggs had progressed beyond MII. In addition, no WT SR-BI+/+ eggs progressed beyond MII at ~16 h (n = 8), indicating that the control SR-BI+/+ eggs may exhibit a mild defect in MII arrest. In other experiments, we observed no differences in the very low level of spontaneous activation between WT and heterozygous mutant eggs. Table 1 shows a dramatic time-dependent increase in SR-BI KO eggs that had progressed past MII at ~16 (73%) or ~20 h (93%) (representative images are in Fig. 3).

Activation of ovulated eggs by either fertilization or parthenogenetic agents suppresses the kinase activities of MPF and MAPK that are required to maintain MII arrest (16, 17, 23). We determined the MPF and MAPK activities of lysates from ovulated WT and SR-BI KO eggs and compared them with those of WT and SR-BI KO oocytes. Table 1 shows a dramatic time-dependent increase in SR-BI KO eggs that had progressed past MII at ~16 (73%) or ~20 h (93%) (representative images are in Fig. 3). Activation of ovulated eggs by either fertilization or parthenogenetic agents suppresses the kinase activities of MPF and MAPK that are required to maintain MII arrest (16, 17, 23). We determined the MPF and MAPK activities of lysates from ovulated WT and SR-BI KO eggs and compared them with those of WT and SR-BI KO oocytes. Table 1 shows a dramatic time-dependent increase in SR-BI KO eggs that had progressed past MII at ~16 (73%) or ~20 h (93%) (representative images are in Fig. 3).

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**Table 1. Quantitative morphologic analysis of spontaneous meiotic progression beyond MII arrest**

<table>
<thead>
<tr>
<th>Stage in meiotic progression</th>
<th>~16</th>
<th>~20</th>
</tr>
</thead>
<tbody>
<tr>
<td>At MII or earlier (%)</td>
<td>93</td>
<td>89</td>
</tr>
<tr>
<td>Beyond MII (%)</td>
<td>7</td>
<td>11</td>
</tr>
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</table>

Four SR-BI KO oocytes at ~16 h and two SR-BI KO oocytes at ~20 h could not be scored, because the chromatin was too diffuse to reliably count centromeres.
activities, impairment of cell cycle control may, in part, account for SR-BI KO female infertility, although other factors may contribute as well.

**Abnormally High Cholesterol Deposition in SR-BI KO EGGS.** In SR-BI KO mice, high cholesterol deposition in RBCs and platelets, caused at least in part by the high UC content of HDL, contributes to diverse abnormalities of these cells (5, 9, 10, 12, 13). We suspected that SR-BI KO eggs might have abnormally high cholesterol accumulation that might contribute to the spontaneous escape of ovulated SR-BI KO eggs from MII arrest. Thus, we measured the relative UC levels in ovulated eggs from normal mice by fluorescence microscopic analysis of cells stained with the cholesterol-binding fluorescent dye filipin (34). Fig. 4A shows that filipin fluorescence in control eggs from SR-BI+/+ females exhibited a weak, uniform intensity together with somewhat brighter puncta. Similar results were observed using eggs from SR-BI−/− females. Eggs from SR-BI+/- females (Fig. 4B) showed a similar staining pattern, except that the intensity of fluorescence was much stronger (68% ± 15% increase, P = 0.004) (Fig. S4). These results indicate the presence of abnormally high levels of UC in the SR-BI KO eggs and raise the possibility that excess UC in SR-BI KO eggs might be responsible for their abnormal escape from MII arrest and possibly might contribute to premature death. Loading cells with excess UC by various routes can induce endoplasmic reticulum (ER) stress, mitochondrial dysfunction, and mitochondrial-mediated cell death (35–41).

**Effects of Excess Cholesterol Loading on WT EGGS.** We directly assessed the ability to activate eggs from WT mice by loading them with excess cholesterol. Exogenous cholesterol was delivered by treating eggs for 10–45 min with 0.125 or 0.5 mM cholesterol-loaded M/C-D (+ cholesterol) as indicated, a standard approach for increasing cellular cholesterol content (26, 42). For example, incubating SR-BI+/+ eggs for 10 min at 37 °C with 0.5 mM cholesterol/M/C-D increased relative UC by 38% ± 10% (P = 0.019) (Fig. S4). Eggs isolated from the oviducts of hormone-primed WT C57BL/6 females ~14–16 h after hCG administration were incubated with or without cholesterol/M/C-D, washed to remove the cholesterol/M/C-D, and then, incubated without cholesterol/M/C-D for an additional 5–6 h at 37 °C. For comparison, some eggs were incubated for 6 h with the well-established chemical activator 5 mM SrCl2 in calcium-free or 2 mM EGTA-containing medium.

We examined several phenotypes associated with egg activation, including (i) alterations in [Ca2+], measured immediately after cholesterol/M/C-D addition (assessed using the Ca2+-sensitive dye Fura-2AM), (ii) MPF and MAPK activities, (iii) second PB extrusion, (iv) stage in meiosis (deconvolution microscopy), and (v) subsequent cleavage to the two-cell stage. Fig. 5 shows typical time courses of the changes in the relative [Ca2+]i after cholesterol addition for five individual WT C57BL/6 eggs continuously incubated with cholesterol/M/C-D for 45 min. For all of the eggs that remained viable during the experiments (n = 26), the exposure to cholesterol/M/C-D induced a single relatively steep increase in [Ca2+]i after an ~15– to 34-min lag. For most of these eggs (24 of 26), the increase was transient, with a partial (3 of 24) or essentially complete (21 of 24) (Fig. 5) return to the baseline before the end of the 45-min imaging period. There was variation in the amplitude and width of the spikes in different eggs. The single spike in [Ca2+]i was somewhat reminiscent of that seen when transient ethanol exposure induces activation (24, 25) and differed from the robust oscillations observed by SrCl2 or fertilization-induced activation (18, 19, 21, 22, 43) (Fig. S5). We also performed [Ca2+]i imaging experiments on eggs exposed to cholesterol/M/C-D for 25 min, in which we imaged the eggs during cholesterol/M/C-D exposure and then, after the cholesterol/M/C-D was washed away. We observed a single large spike in [Ca2+]i that returned to baseline when we collected data between 20 and 155 min after removing the cholesterol/M/C-D (Fig. S6). There was no evidence of [Ca2+]i oscillations induced by cholesterol/M/C-D.

Fig. 6 shows that cholesterol loading for 20–25 min seemed to be as effective as SrCl2 treatment in activating the eggs, which was assessed by suppression of MAPK and MPF kinase activities (Fig. 6A), induction of PB extrusion (Fig. 6B and C), and progression to subsequent stages in meiosis (e.g., prometaphase formation) (Fig. 6C). All three assessments were made ~5–6 h after the addition of cholesterol or SrCl2. Comparable cholesterol induction of second PB extrusion was observed using eggs from hormone-primed SR-BI+/- mice on a mixed C57BL/6:129 background (Fig. S7). We also measured second PB extrusion in WT C57BL/6 eggs during [Ca2+]i imaging experiments, in which we imaged the eggs during the 25-min cholesterol/M/C-D exposure and for an additional 15 min after the cholesterol/M/C-D was washed away. A single [Ca2+]i spike was observed in most of the
viable eggs (16 of 18 in three independent experiments) (Fig. S6B), and 94% (15 of 16) of those eggs extruded a second PB by the end of the second 155-min cholesterol/Müller-free incubation. Thus, multiple distinct [Ca^{2+}]” spikes were not required for cholesterol-induced second PB extrusion.

In multiple experiments, we found that most cholesterol/Müller-activated eggs progressed to the two-cell stage 24 h after treatment. For example, in two such experiments, we followed 24 h in vitro the fates of 0.125 (n = 41) or 0.5 mM (n = 11) cholesterol-loaded WT C57BL/6 eggs; of those eggs that extruded second PBs after 6 h (68% and 73%, respectively), 71% and 63%, respectively, underwent cleavage to the two-cell stage by 24 h after cholesterol loading. In contrast, untreated eggs remained arrested in MIi. Strikingly, in our [Ca^{2+}]” imaging experiments on C57BL/6 eggs loaded with cholesterol for 25 min, of those eggs that were viable 24 h after treatment (n = 29), 90% extruded a second PB at ~6 h (n = 26), 100% of which exhibited a Ca^{2+} spike. In addition, 86% of those activated eggs that were examined (19 of 22) were found to have progressed to the two-cell stage. Of the remaining three eggs that had not extruded a second PB, two eggs exhibited a Ca^{2+} spike, and none progressed to the two-cell stage (additional details in SI Materials and Methods). Thus, induction of a Ca^{2+} spike may be necessary but not sufficient for cholesterol/Müller-activated egg activation. Future experiments will be required to determine which features of the Ca^{2+} spike (onset time, duration, amplitude, etc.) and the status of the eggs (e.g., extent of cholesterol loading) are necessary for activation. Taken together, our observations indicate that delivery of exogenous cholesterol to eggs efficiently induces exit from MIi arrest by mechanisms that include biochemical changes associated with classic chemical-mediated activation and sperm-mediated activation during fertilization. Thus, cholesterol delivery can induce activation, suggesting that spontaneous activation caused by excess cholesterol in eggs of SR-BI KO females contributes directly to their female infertility.

Discussion

Female SR-BI KO mice exhibit fully penetrant infertility (6, 14). Our previous findings that these animals have essentially normal ovarian histology and that their fertility can be readily rescued by an HDL cholesterol-lowering drug suggested that the infertility might be a consequence of defects in oocytes or eggs, possibly caused by the animals’ unusual dyslipidemia (hypercholesterolemia with abnormal HDL structure and composition) (2, 6, 7, 14, 15). Indeed, defects occurring during or soon after ovulation in vivo are readily apparent, because ~19% of the eggs harvested ~16 h after hCG administration (hormone-induced superovulation) are dead (refs. 6 and 14 and this study).

Here, we sought to understand the cellular basis of SR-BI KO female infertility by investigating oocyte growth and meiotic progression. We observed no obvious defects in oocyte growth in the follicle, maintenance of primary meiotic arrest in prophase I, resumption of meiosis after release from primary arrest (kinetics of GVBD), or efficiency of first PB extrusion (Fig. 1 shows a schematic depiction of meiosis and parthenogenesis). However, when we examined ovulated eggs from control and SR-BI KO females, we found that substantial numbers of SR-BI KO eggs had spontaneously activated (escaped MIi arrest and exhibited second PB extrusion and suppression of MFP and MAPK activities). Deconvolution immunofluorescence imaging of the untreated, viable SR-BI KO eggs harvested at ~16 or ~20 h after hCG administration established that the progressive spontaneous escape from MIi arrest resulted in second PB extrusion, pronucleus formation, and progression to metaphase III and

Fig. 6. Additional effects of cholesterol loading and SrCl2 treatment on eggs from WT C57BL/6 females. (A–C) Eggs from WT C57BL/6 females were harvested ~14–16 h after hCG administration and stripped of cumulus cells by hyaluronidase treatment. (A) MAPK and MPF kinase activities. Eggs were pretreated for 25 min without (−) or with (+) 0.125 mM cholesterol-loaded MÜller (Cholesterol) in M16 at 37 °C as indicated. They were then washed and incubated in M16 medium either without (untreated (n = 60) or cholesterol-loaded MÜller (n = 68) eggs) or with 5 mM SrCl2/2 mM EGTA (n = 40) at 37 °C for 6 h. Eggs were then washed, pooled into sets of four, and lysed, and MAPK and MPF kinase activities were measured as described in SI Materials and Methods. The values represent the averages ± SEMs from four independent assays using eggs harvested and processed on 7 different d. The values for one-way ANOVA for MPF and MAPK activities were P < 0.00001. *P value < 0.02 (unpaired Student t test) for comparisons with SrCl2. †P value < 0.0001 (unpaired Student t test) for comparisons with untreated controls; ‡P value < 0.001 (unpaired Student t test) for comparisons with SrCl2. (B) PB extrusion. Some of the eggs (n = 124) were pretreated at 37 °C for 25 min with (+) or without (−) 0.125 mM cholesterol-loaded MÜller (Cholesterol) in M16 medium and then washed and incubated in M16 medium for an additional 6 h at 37 °C. The remaining eggs were incubated for 6 h at 37 °C in M16 medium without (n = 103) or with (n = 114) 5 mM SrCl2/2 mM EGTA. The fraction of eggs exhibiting second PB extrusion was then determined (averages from eight independent experiments). The value for one-way ANOVA was P < 0.00001. *P value < 0.00001 (paired Student t test) for comparisons with untreated controls. (C) Deconvolution microscopy. Eggs were pretreated for 20 min without (−) or with (+) 0.5 mM cholesterol-loaded MÜller (Cholesterol) in M16 at 37 °C as indicated. They were then washed and incubated in M16 medium either without (untreated (n = 20) or cholesterol-loaded (n = 20) eggs) or with 5 mM SrCl2/2 mM EGTA (n = 17) at 37 °C for 6 h. We then stained the cells and collected deconvolved and z-projected images as described in Fig. 3. (Upper) Low-magnification images show the overall morphology of the eggs; dashed lines outline the cells and the second PBs. (Lower) High-magnification images show the number and arrangement of chromatids in either the main cell [untreated; 40 paired centromeres (red) at the MIi spindle] or the second PBs (SrCl2 and cholesterol). The fractions of eggs that extruded a second PB were untreated, 0% (0 of 18); SrCl2, 77% (10 of 13); and cholesterol/MÜller, 74% (14 of 19). Those eggs that were fragmented at the end of the 6-h incubation (untreated, two; SrCl2, four; cholesterol, one) were not included in the calculation of the percentage exhibiting second PB exclusion. ACA, anticentromere antibody. (Scale bar: 10 μm.)
anaphase/telophase III. Spontaneous fertilization-independent exit from MII (or inability to establish or maintain arrest at MII) results in cells with a haploid chromosome complement, which in at least some cases, is further reduced during subsequent progression to metaphase III and anaphase/telophase III. These cellular defects are likely to account for at least some and perhaps most of the inability of ovulated SR-BI eggs to be fertilized and develop normally into viable embryos after mating (6). Additional studies are necessary to determine whether the SR-BI KO eggs that were dead at harvesting ~16 h posthormone induction died because of meiotic defects or if other independent abnormalities also contributed to this process.

**Cholesterol Accumulation in SR-BI KO Eggs Contributes to Their Abnormal Phenotypes.** One potential mechanism underlying premature death and spontaneous escape from MII arrest of SR-BI KO eggs is their abnormal accumulation of cholesterol. Indeed, excess cholesterol accumulation in RBCs and platelets in SR-BI KO mice, apparently caused by the high UC:TC ratio in circulating HDL (7, 8), has previously been associated with the abnormal maturation, lifetime, structure, and/or function of these blood cells (7–13). As was the case for RBCs and platelets in SR-BI KO mice, we detected excess cholesterol in ovulated SR-BI KO eggs (here assessed by filipin staining). In complementary experiments, we found that cholesterol loading of control, fertile, SR-BI/−/−, or WT eggs by incubation with a cholesterol/MjCD complex resulted in activation similar to that of SR-BI KO eggs as well as a single spike in [Ca\(^{2+}\)]. Prolonged accumulation of excess cholesterol might also contribute through various routes (e.g., ER stress, mitochondrial dysfunction, and excessive or prolonged [Ca\(^{2+}\)] elevation) to the premature death of some of the SR-BI KO eggs (35–41, 44, 45). We, therefore, conclude that (i) cholesterol can induce parthenogenetic activation of mouse eggs and that (ii) accumulation of excess cholesterol in SR-BI KO oocytes plays a substantial role in generating their abnormal phenotypes and consequently, the infertility of SR-BI KO females. Conversely, Buschiazzo et al. (46) have shown that cholesterol depletion by exposure to high levels of cholesterol-free MjCD (15 mM for 30 min at 37 °C) induces a delay in second PB extrusion and a low fertilization of mouse eggs. We have found that pretreating SR-BI/−/− and SR-BI KO eggs with 10 mM cholesterol-free MjCD reduces filipin staining intensity by >80% and reduces fertilization rate by >88% (reduced viability, respectively). Taken together, these data suggest that cholesterol in mouse eggs must be maintained within a relatively narrow physiological range to ensure the oocyte’s health and developmental potential. A critical role of UC in mediating oocyte/egg abnormalities is consistent with our previous report that correcting the abnormal composition (high UC:TC ratio) of plasma HDL can restore fertility (7, 10, 14). It is possible that excess cholesterol-independent mechanisms may also be involved.

**HDL Is the Likely Source of Excess Cholesterol in SR-BI KO Eggs.** Four lines of surgical transplantation, genetic, and pharmacologic evidence support the hypothesis that abnormal circulating HDL in SR-BI KO mice (high UC:TC ratio and large diameter) (5, 7, 8) is a major cause of female SR-BI KO infertility and likely to be the cause of excess cholesterol accumulation in their eggs. (i) Ovary transplantation established that ovarian expression of SR-BI is not necessary for fertility during embryonic development (primordial follicle production) or adult maturation and function (primordial follicle conversion to antral follicles and ovulation). SR-BI KO ovaries bilaterally transplanted into SR-BI-positive hosts developed the capacity to ovulate functional eggs that can be fertilized and generate healthy pups (14). (ii) Tissue-specific transgenic expression of SR-BI in the liver of SR-BI KO mice restores not only normal HDL composition and structure but also, essentially normal fertility (15). (iii) In SR-BI KO mice, homozygous genetic ablation of the gene encoding apoA-I, the major protein component of HDL, reduces total plasma cholesterol (mainly HDL cholesterol) to essentially WT levels, although the HDL particles remain abnormally large. Fertility is partially restored in these double KO mice (41% fertility and small litter sizes) (14). (iv) Oral administration of the HDL cholesterol-lowering drug probucol reduces total plasma cholesterol to WT levels and restores to essentially normal the UC content of HDL, although the HDL particles remain abnormally large (7, 14). Fertility of SR-BI KO females is restored soon after probucol treatment (14). Several studies indicate that the abnormal UC composition (high UC:TC ratio) rather than the increased size of HDL particles in SR-BI KO mice may play a particularly important role in SR-BI KO female infertility. As noted above, apoA-I deficiency and probucol treatment partially or nearly fully restore fertility to SR-BI KO mice without substantially reducing the abnormally large size of HDL particles. Also, transgenic expression of the human cholesterol ester transfer protein in SR-BI KO mice reduces the size/density of the HDL in a dose-dependent fashion (47, 48). Although the size of the HDL particles in these transgenic SR-BI KO mice is normalized, the composition is not (abnormally high UC:TC ratio), and they are infertile (48).

There are two attractive mechanisms by which abnormal HDL particles in the follicular fluid of antral follicles in SR-BI KO mice might lead to excess cholesterol accumulation in oocytes. Because of the relatively small size of HDL particles compared with other lipoproteins, HDL in many reproductive tissues may contain different lipids in substantially greater amounts in the follicular fluid that bathes the developing oocyte in the antral follicle (49–52). It may be that, in the follicle, the UC-enriched HDL particles transfer cholesterol to the oocytes directly, which they seem to do in RBCs and platelets (9, 10), or indirectly (e.g., through surrounding cumulus cells). Su et al. (53) have suggested that mouse preovulatory oocytes synthesize little cholesterol and obtain cholesterol primarily from the surrounding cumulus cells. Thus, there seems to be a mechanism that could transfer HDL-derived cholesterol from cumulus cells to the oocytes. An alternative for excess cholesterol accumulation in SR-BI KO oocytes is that normal cholesterol homeostasis in oocytes may depend on adequate levels of HDL-mediated cholesterol efflux to balance input and that the UC-enriched HDL particles in SR-BI KO mice may be unable to mediate sufficient cholesterol efflux to prevent excess cholesterol accumulation.

HDL has been shown to have multiple activities, including the ability to transport lipids other than cholesterol [e.g., vitamin E (54) and sphingosine-1-phosphate (55)] and influence oxidation, inflammation, metabolism, chemotaxis, thrombosis, microbial infection, cell signaling, and others (56–58). Thus, it is possible that, in addition to contributing to accumulation of excess cholesterol in oocytes, some of the effects of the abnormal HDL on female fertility in SR-BI KO mice may be independent of its role in cholesterol transport.

**Excess Cholesterol and Egg Activation.** One possible mechanism by which excess cholesterol might activate eggs is its stimulation of an increase in cytoplasmic [Ca\(^{2+}\)], possibly by inducing ER Ca\(^{2+}\) release through a stress pathway or by otherwise influencing calcium channel or pump activities. Normally, fertilization or parthenogenetic agents (e.g., SrCl\(_2\)/EGTA or ethanol) induce a single spike or oscillations in [Ca\(^{2+}\)], that are important for activation and successful progression of embryonic development (18, 19, 24, 25, 43).

Excess cholesterol-induced escape from MII arrest is unlikely to involve all of the same mechanisms as those induced by the MEK inhibitor U0126 (blocks MAPK activation) (23, 59, 60), c-mos deficiency (23, 61), or Zn depletion (25, 62, 63) because of their differing phenotypes. Both c-mos KO and Zn-insufficient eggs share with SR-BI KO eggs an otherwise spontaneous exit from MII arrest (64). However, unlike SR-BI KO eggs, c-mos KO and
Zn-insufficient eggs often extrude abnormally large first PBs (64). Aged eggs—those not fertilized within a critical time after ovulation (65–68)—also are subject to relatively high rates of spontaneous activation that involves exit from MII arrest, second PB extrusion, arrest at a metaphase III-like stage (abnormal spindle and no pronucleus), and cell death (fragmentation and apoptosis) (68, 69). Future studies will be required to characterize in detail the mechanisms underlying excess cholesterol-induced activation. Those future studies will help determine if cholesterol-induced activation might serve as an alternative to SrCl2-induced or other (70) activation methods and perhaps, even improve efficiency in procedures, such as somatic cell nuclear transfer, to derive embryonic stem cells from nuclear transfer blastocysts or generate cloned animals (70, 71).

Implications for Human Infertility. This study has uncovered a previously unidentified ability of excess cholesterol to activate MII-arrested mouse eggs. In addition, this report raises the possibility that abnormalities in cholesterol metabolism in general and SR-BI–mediated cholesterol metabolism specifically (6, 7, 14) have the potential to contribute to human female infertility of unknown etiology (72, 73). In such cases, pharmacologic manipulation of cholesterol metabolism (e.g., lowering excess cholesterol in oocytes or eggs) might prove beneficial. It is noteworthy that Velasco et al. (74) and Yates et al. (75) have reported a potential influence of variations in SR-BI on fertility of women undergoing in vitro fertilization (IVF). In one study, Velasco et al. (74) found that SR-BI levels in granulosa cells from 50 volunteers were correlated with plasma levels of estradiol and the number of retrieved and fertilized oocytes, although SR-BI levels were not correlated with quality measurements of embryos, primary diagnosis of infertility, or pregnancy outcomes. A study of 274 women by Yates et al. (75) found no association of the SR-BI alleles tested with qualitative measures of embryo viability (blastocyst number and grade). However, 0 of 10 heterozygous carriers of one minor allele (rs4238001) had viable fetuses after embryo transfer, raising the possibility that this minor allele (and another allele in an African-American subset of subjects) may possibly contribute to human female infertility. A recent report suggests that there may be an association of high-plasma UC in males and females and increased time to pregnancy in 501 couples (76). In addition, the report of a case of human infertility caused by spontaneous activation (77) raises the possibility that excess cholesterol-induced egg activation may have clinical relevance.

Despite the progress in assisted reproduction in the last decades, infertility remains an unresolved medical problem. Recent data indicate that there are more than 30 million women with infertility of unknown origin (78). Additional studies will determine if and to what extent variations in SR-BI activity specifically or cholesterol deposition in oocytes or eggs generally might influence human female fertility.

Materials and Methods

Pure C57BL/6 (Charles River Laboratories, Jackson Laboratory, and Taconic), SR-BI+/-, SR-BI-/-, and SR-BI+/- (S) mice were maintained on an RMH3000 laboratory diet ad libitum, and experiments were performed in concordance with the guidelines from the committees for animal care at the Massachusetts Institute of Technology and Pontificia Universidad Católica de Chile (SI Materials and Methods). Oocytes were collected from unprimed or primed females, oocyte and follicle diameters were measured, in vitro maturation of oocytes was performed, eggs were activated with SrCl2/EGTA, cholesterol was depleted by MjCD or delivered by cholesterol/MjCD complexes, relative cholesterol content was assessed using filipin staining, eggs were imaged by whole-mount staining and deconvolution microscopy, and statistical analyses were performed using standard procedures (SI Materials and Methods).

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