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Steroid receptor coactivators 1 and 2 mediate fetal-to-maternal signaling that initiates parturition

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Introduction

Preterm birth is a leading cause of neonatal morbidity and mortality worldwide (1–4). Globally, approximately 15 million babies are born prematurely each year (1) and approximately 1.1 million babies die annually from complications of prematurity (4). The initiation of labor, both at term and preterm, is associated with an upregulated inflammatory response, exemplified by increased proinflammatory cytokines in amniotic fluid (AF) (5); infiltration of myometrium, cervix, and fetal membranes by neutrophils and macrophages (6–8); and a decline in progesterone (P₄) receptor (PR) function. In preterm labor, intraamniotic (i.a.) infection likely provides a prominent stimulus for increased AF cytokines and inflammatory cell migration (9, 10). However, the precise signals for the inflammatory response leading to labor at term remain unclear (11).

Parturition timing in all species appears to be mediated by signals from both mother and fetus (11). Near term, increased uterine stretch (12, 13) and factors produced by the developing fetus (6, 14–16) may provide inflammatory stimuli. Previously, we (6) and others (17) suggested that the fetus may provide a signal via pulmonary surfactant, a developmentally regulated, glycerophospholipid-rich lipoprotein produced by pulmonary alveolar type II cells and secreted into AF in increased amounts during late gestation (11). Surfactant, which reduces surface tension within the lung alveoli after birth, is essential for air breathing. Thus, infants born prematurely are at risk of developing respiratory distress syndrome because of surfactant deficiency. We further suggested that the major surfactant protein, surfactant protein-A (SP-A), a C-type lectin also secreted by the fetal lungs into AF near term, could serve as a hormonal signal for parturition (6). SP-A stimulated IL-1β expression in cultured AF macrophages, and i.a. injection of SP-A caused preterm delivery of fetuses within 6 to 24 hours (6). In mice, targeted deletion of the genes encoding SP-A and the related C-type lectin, SP-D, caused a modest but significant parturition delay and prevented induction of “contractile” genes, oxytocin receptor (Oxtr) and connexin-43 (Cx43), also known as gap junction protein 1 (Gja1), in myometrium (15). While these findings were compelling, both parents were double KO (dKO) for SP-A and SP-D. Thus, a role for fetal signaling was not definitively proven. Moreover, the parturition delay was observed only in the second or subsequent pregnancies, suggesting a requirement for other signals.

Our previous studies revealed that a 300-bp region upstream of the SP-A gene mediates its developmental, tissue-specific, and
hormonal regulation of expression (18, 19). Steroid receptor coactivators 1 and 2, SRC-1 and SRC-2, serve important roles in transcriptional upregulation of SP-A gene expression in fetal lung type II cells (20, 21). Endogenous SRC-1 in human fetal lung (HFL) type II cells was recruited with thyroid transcription factor-1 (TTF-1, also known as Nkx2.1) and NF-κB to the hSP-A promoter upon cAMP and IL-1 stimulation of SP-A expression (21, 22). cAMP also enhanced estrogen-related receptor α (ERRα) transcriptional activity at the SP-A promoter by increasing its interaction with protein kinase A and SRC-2 (20).

SRCs belong to the p160 coactivator family, comprised of three homologous members: SRC-1 (also known as ERAP140, ERAP160, and NCOA1), SRC-2 (also known as TIF-2, GRIP-1, and NCOA2), and SRC-3 (also known as p/CIP, RAC3, AIB1, TRAM-1, and ACTR). Although they do not directly bind to DNA, coactivators can robustly increase the transcriptional activities of nuclear receptors and other transcription factors by bridging them to factors bound to other response elements and to the basal transcription machinery and by recruiting other coregulators (23, 24). Through recruitment of coregulators causing posttranslational modification of histones, they can alter chromatin structure, resulting in transcriptional activation or repression (25).

Studies using KO mice indicate that, although structurally similar, each SRC family member has specific physiological functions (24). For example, although SRC-1/- mice are fertile, female mice manifest reduced estrogen-stimulated uterine growth and decreased P4-dependent uterine decidual response (26). Male SRC-2/- mice are infertile due to defects in spermatogenesis (27), while female SRC-2/- mice display placental hypoinsulinemia and hypo-fertility (28). In light of our previous findings that SRC-1 and SRC-2 serve roles in the regulation of SP-A gene expression, we were intrigued by the finding that mice lacking both SRC-1 and SRC-2 (SRC-1/- SRC-2/- mice, herein referred to as SRC-1/- dKO mice) died at birth of respiratory distress (29), suggesting a deficiency of pulmonary surfactant production.

To investigate the molecular basis for effects of SRC-1 and SRC-2 on surfactant synthesis by the fetal lung, we analyzed SRC-1 and SRC-2 double-deficient mice. Unexpectedly, we uncovered a pathway that we believe to be novel by which SRC-1 and SRC-2 coordinately control pregnancy length and the initiation of labor through their impact on production of surfactant lipid and proteins by the fetal lung.

Results

SRC-1/SRC-2 double deficiency in mice results in delayed labor. To define the phenotypic consequences of deficiency in SRC-1 and SRC-2, we crossed male and female mice that were double heterozygous for SRC-1 and SRC-2 (SRC-1/- SRC-2/- mice, herein referred to as SRC-1/- dhet mice). To ensure that parturition timing was analogous across all genotypes to be similar (Supplemental Table 1; supplemental material available online with this article; doi:10.1172/JCI78544DS1). Notably, SRC-1/- dhet females bred to genotypically matched males or WT females bred to 1-KO/2-het males manifested a similar delayed labor during second and subsequent pregnancies (Supplemental Table 2).

The litters generated from SRC-1/- dhet female mice bred to SRC-1/- dhet male mice are expected to be comprised of 9 possible genotypes according to Mendelian law (Figure 1A). Upon genotyping all of the fetuses in each litter, we plotted the proportion of SRC-1 SRC-2 double-deficient pups (which included SRC-1/- dhet, SRC-1/- dKO, 1-KO/2-het, SRC-1/- SRC-2/- pups [herein referred to as 1-het/2-KO pups]) for each of the crosses relative to gestational length. A strongly positive correlation was found between the proportion of SRC-1 SRC-2 double-deficient pups in corresponding litters and the timing of parturition (Figure 1C).

SP-A synthesis and secretion are impaired in fetal lungs of SRC-1/- double-deficient mice. In light of the evidence that the delayed-labor phenotype was caused by an alteration in signals produced by SRC-1/- double-deficient fetuses, as well as our previous findings that SRC-1 and SRC-2 are important for transcriptional regulation of SP-A in fetal lung type II cells (20, 21), we analyzed the SP-A mRNA and protein expression in fetal lungs and secretion of SP-A protein into AF. Notably, SP-A mRNA and protein in lungs of SRC-1/- double-deficient fetuses were significantly decreased compared with those of WT fetuses and with those of fetuses that were singly deficient in SRC-1 or SRC-2 (Figure 1, D and E). Moreover, SP-A protein in AF of SRC-1/- double-deficient fetuses also was dramatically decreased compared with that of WT fetuses and with that of fetuses that were singly deficient for either coactivator (Figure 1F).

To determine whether other surfactant proteins were affected by SRC-1/- deficiency, we analyzed expression of SP-D, a C-type lectin structurally similar to SP-A, as well as the hydrophobic surfactant protein, SP-B, which is essential for air breathing (30, 31). Expression of these two surfactant proteins was unaffected in lung tissues of SRC-1/- double-deficient fetal mice (Supplemental Figure 1, A and B). Additionally, the ATP-binding cassette, subfamily A, member 3 (ABCA3), an essential transporter that mediates surfactant lipoprotein trafficking and lamellar body biogenesis (32, 33), showed only modest, nonsignificant changes in lungs of SRC-1/- double-deficient fetuses (Supplemental Figure 1C).
To confirm the KO efficiencies of SRC-1 and SRC-2 in fetal lungs, we performed real-time PCR and found that SRC-1 and SRC-2 mRNA were essentially undetectable in SRC-1 KO fetal lungs and SRC-2 KO fetal lungs, respectively. Neither was detected in SRC-1/-2 dKO fetal lungs. Interestingly, SRC-1 mRNA expression was markedly and significantly increased in fetal lungs of SRC-2 KO mice, while SRC-2 mRNA was significantly increased in fetal lungs of SRC-1 KO mice (Supplemental Figure 1, D and E, respectively), suggesting that SRC-1 and SRC-2 are upregulated in a compensatory manner in vivo.

NF-κB activation is decreased in myometrium of WT or SRC-1/-2-deficient females bred to SRC-1/-2-deficient males. We previously suggested that the developmental increase in SP-A expression in mouse fetal lung and its secretion into the AF after 17 dpc can activate AF macrophages via binding to TLR2. The activated macrophages, in turn, may transit to the maternal uterus, resulting in increased cytokine production and myometrial NF-κB activation (6, 15). To determine whether a deficiency of SRC-1/-2 affected the activation of inflammatory signaling pathways, the proinflammatory transcription factors, NF-κB p65 and p50, were analyzed in myometrial cytoplasmic and nuclear fractions by immunoblotting. Notably, nuclear levels of NF-κB p65 and p50 proteins were greatly reduced in myometrial tissues of SRC-1/-2 dko mice bred to SRC-1/-2 double-deficient males relative to WT females bred to SRC-1/-2 double-deficient males at 18.5 dpc, in association with a coordinate increase in cytoplasmic levels of these proteins (Figure 2, A and B). These findings suggest that inflammatory stimuli promoting nuclear translocation and activation of NF-κB were impaired in the pregnant uterus near term, in association with the decrease in SP-A expression and secretion in SRC-1/-2-deficient fetal mice.

Expression of contractile genes is decreased in myometrium of WT or SRC-1/-2-deficient females bred to SRC-1/-2-deficient males. The contractile genes Cx43, Oxtr, and cyclooxygenase 2 (Cox2,
Figure 2. NF-κB activation and contraction-associated protein gene expression are decreased in 18.5 dpc pregnant myometrium of SRC-1/-2 dhet females × SRC-1/-2 dhet males and WT females × 1-KO/2-het males, compared with WT females × WT males. (A and B) Nuclear translocation of (A) NF-κB p65 and (B) p50 in myometrium. Representative immunoblots of p65 and p50 protein in nuclear (Nuc) or cytoplasmic (Cyto) fractions. The same immunoblots of histone H3 and GAPDH are shown for nuclear and cytoplasmic proteins, respectively, since p65 and p50 were probed on the same respective blots. Combined data from densitometric scans of nuclear immunoblots for p65 and p50 relative to cytoplasmic immunoblots, normalized to internal controls for WT females × WT males (n = 10), WT females × 1-KO/2-het males (n = 8), and dhet females × dhet males (n = 10). (C) Cx43 mRNA and (D) protein levels for WT females × WT males (n = 5), WT females × 1-KO/2-het males (n = 4), and dhet females × dhet males (n = 5). The same immunoblot of β-actin (D) was used for the normalization of AKR1B3 (Figure 3B) and PGDH (Figure 3D), which were probed on same blot. (E) Oxtr mRNA and (F) protein in myometrium from WT females × WT males (n = 5), WT females × 1-KO/2-het males (n = 4), and dhet females × dhet males (n = 5). Densitometric scans were normalized to β-actin. Data are mean ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001 (ANOVA).
also known as prostaglandin-endoperoxide synthase 2 \( [\text{Ptgs2}] \), the critical enzyme in synthesis of contractile prostaglandins, which promote transformation of the myometrium from a quiescent state to a contractile unit, are known targets of NF-kB (34, 35). To determine the direct cause of delayed initiation of myometrial contraction in \( \text{SRC-1/-2} \)-deficient mice, the expression of these contractile genes in myometrium was analyzed. mRNA and protein levels of both CX43 and OXTR were significantly decreased in myometrium of \( \text{SRC-1/-2} \) dhet or WT females bred to 1-KO/2-het males compared with those in WT crosses (Figure 2, C-F), whereas no significant differences in \( \text{Cox2} \) mRNA or protein levels were detected among these groups of mice (Supplemental Figure 2, A and B). Interestingly, expression of the prostaglandin \( F_\alpha \)-specific (PGF\( _\alpha \)-specific) synthase, AKR1B3, was dramatically decreased, while that of the PGF\( _\alpha \)-metabolizing enzyme, prostaglandin dehydrogenase (PGDH), was significantly increased in 18.5-dpc myometrium of \( \text{SRC-1/-2} \) dhet females bred to genetically matched males or WT females bred to 1-KO/2-het males (Figure 3, A-D). These changes were associated with a significant decrease in myometrial PGF\( _\alpha \) levels (Figure 3E).

**Pregnant **\( \text{SRC-1/-2} \)-deficient or WT females bred to \( \text{SRC-1/-2} \)-deficient males manifest delayed luteolysis, evidenced by increased ovarian \( \text{StAR} \) expression and serum \( P_4 \) levels at term. In a number of mammalian species, uterus-derived PGF\( _\alpha \) plays an important role in controlling luteolysis (36). In late gestation, activation of inflammatory response pathways leads to upregulation of uterine PGF\( _\alpha \), which then circulates to the ovary, in which it binds to PGF\( _\alpha \) receptors to initiate corpus luteum regression (36, 37). In rodents, \( P_4 \) production by the ovarian corpus luteum is maintained at high levels throughout pregnancy and diminishes near term, due to luteolysis. Increased \( P_4 \) production is maintained, in part, by steroidogenic acute regulatory protein (\( \text{StAR} \)), which facilitates translocation of cholesterol from the outer to the inner mitochondrial membrane, in which cholesterol is metabolized to pregnenolone by cholesterol side-chain cleavage (P450scc) (38, 39), the rate-determining step in steroidogenesis. \( \text{StAR} \) expression declines in the corpus luteum near term, with an associated fall in \( P_4 \) production (36). In this study, we found that circulating \( P_4 \) levels (Figure 3F) and ovarian \( \text{StAR} \) protein expression (Figure 3G) failed to decrease in \( \text{SRC-1/-2} \) dhet or WT females bred to \( \text{SRC-1/-2} \) dhet or 1-KO/2-het males, respectively. Using immunohistochemical analysis, we found prolonged \( \text{StAR} \) expression in the corpus luteum of the \( \text{SRC-1/-2} \) dhet females bred to \( \text{SRC-1/-2} \) dhet males (Figure 3H). This was clearly in contrast to WT mice, in which a pronounced decline in \( \text{StAR} \) expression occurred in late gestation (Figure 3H). Interestingly, \( \text{StAR} \) mRNA expression was not significantly altered in ovaries of WT or \( \text{SRC-1/-2} \) dhet females bred to \( \text{SRC-1/-2} \) dhet or 1-KO/2-het males as compared with that of WT females bred to WT males near term (Supplemental Figure 3A). Similarly, PGF\( _\alpha \) was found to reduce \( \text{StAR} \) protein, but not mRNA, levels in rat luteal cells (40), suggesting posttranscriptional regulation. We also analyzed myometrial expression of the \( P_4 \)-metabolizing enzyme 20α-hydroxysteroid dehydrogenase (20α-HSD), which increases in myometrium near term (41), but failed to observe significant changes in its protein levels with \( \text{SRC-1/-2} \) deficiency (Supplemental Figure 3C). Moreover, we correlated the degree of \( \text{SRC-1/-2} \) double deficiency (by the proportion of \( \text{SRC-1/-2} \) double-deficient fetuses in each litter) with circulating levels of maternal \( P_4 \) (as a measure of luteolysis) and found a strong positive correlation between \( P_4 \) levels and the degree of \( \text{SRC-1/-2} \) double deficiency (Supplemental Figure 3D). These collective findings suggest that fetal \( \text{SRC-1/-2} \) deficiency inhibits a cascade of signals that promote increased PGF\( _\alpha \) production in the uterus and luteolysis at term. This results in maintenance of elevated circulating maternal \( P_4 \) in \( \text{SRC-1/-2} \)-deficient mice, which likely contributes to the delay in parturition.

**LPCAT1 expression and synthesis of platelet-activating factor are reduced in lungs of \( \text{SRC-1/-2} \)-deficient fetuses.** As mentioned, targeted deletion of the genes encoding \( \text{SP-A} \) and \( \text{SP-D} \) in mice also caused a significant delay in parturition (14.4 ± 4.3 hours) (15). However, WT females bred to 1-KO/2-het males manifested a much more striking delay in parturition (38.4 ± 1.0 hours) (Supplemental Figure 4), suggesting the potential roles of fetal signals other than \( \text{SP-A} \) and \( \text{SP-D} \) in the timing of labor. Indeed, the delay in parturition in the \( \text{SRC-1/-2} \)-double-deficient mice may have been underestimated, since those pregnant mice that failed to deliver 22.5 dpc were sacrificed. These differences in gestation length suggest the role(s) of fetal factors other than surfactant proteins in parturition timing.

Previously, it was observed that \( \text{SRC-1/-2} \)-dKO fetuses died at birth of apparent respiratory distress. The lungs of these fetuses manifested alveolar collapse (29), possibly due to impaired surfactant glycerophospholipid synthesis. To determine the enzyme(s) in the surfactant glycerophospholipid pathway that may be affected by the combined deficiency in \( \text{SRC-1/-2} \), we analyzed expression of those enzymes catalyzing the key metabolic reactions. These included CTP:phosphocholine cytidylyltransferase \( \alpha \) (CCT\( \alpha \)), which catalyzes the condensation of CTP and phosphocholine to form CDP-choline; cholinephosphotransferase I (CHPTI), which catalyzes the reaction of CDP-choline and diacylglycerol to form phosphatidycholine; and lysophosphatidylcholine acyltransferase-1 (LPCAT1), a key enzyme in surfactant phosphatidylcholine remodeling to form dipalmitoylphosphatidylcholine (DPPC), the major surface-active component of surfactant (42–44). Notably, neither CCT\( \alpha \) nor CHPTI manifested significant changes in expression in lungs of \( \text{SRC-1/-2} \) double-deficient fetuses compared with WT fetuses (Supplemental Figure 5, A and B). By contrast, \( \text{Lpcat1} \) mRNA and protein levels were markedly increased in WT fetal lungs at 18.5 dpc, compared with 15.5 dpc; however, this developmental increase failed to occur in lungs of \( \text{SRC-1/-2} \) double-deficient fetuses (Figure 4, A and B). This developmental induction of LPCAT1 in WT mice was consistent with a previous report indicating that LPCAT1 was developmentally regulated in the fetal lung and expressed specifically in type II cells (43). Intriguingly, we observed that expression of \( \text{Lpcat1} \) mRNA and protein was significantly decreased in lungs of \( \text{SRC-1/-2} \) double-deficient fetuses compared with that of WT fetuses (Figure 4, C and D). By using liquid chromatography mass spectrometry (LC-MS), we observed that DPPC concentrations were dramatically decreased in AF surrounding \( \text{SRC-1/-2} \) dhet and \( \text{SRC-1/-2} \) dKO fetuses at 18.5 dpc (3.54 ± 1.68 ng/μl and 2.70 ± 0.67 ng/μl, respectively, \( P < 0.0001 \)), compared with WT fetuses (9.05 ± 1.80 ng/μl), while DPPC levels in AF surrounding...
Figure 3. SRC-1/-2 dhet or WT females bred to dhet or 1-KO/2-het males, respectively, manifest decreased myometrial PGF$_2\alpha$ synthesis and delayed ovarian luteolysis. (A) Akr1b3 mRNA and (B) protein in myometrium from WT females × WT males (n = 5), WT females × 1-KO/2-het males (n = 4), and dhet females × dhet males (n = 5). Immunoblots and densitometric scans normalized to β-actin are shown. (C) PGDH mRNA and (D) protein in myometrium of WT females × WT males (n = 5), WT females × 1-KO/2-het males (n = 4), and dhet females × dhet males (n = 5). Immunoblots and densitometric scans normalized to β-actin are shown. (E) PGF$_2\alpha$ levels in myometrial homogenates normalized to protein for WT females × WT males (n = 12), WT females × 1-KO/2-het males (n = 8) and dhet females × dhet males (n = 15). In the box-and-whisker plot, horizontal bars indicate the medians, boxes indicate 25th to 75th percentiles, and whiskers indicate 10th and 90th percentiles. (F) Maternal serum P$_4$, measured by ELISA for WT females × WT males (n = 12), WT females × 1-KO/2-het males (n = 8), and dhet females × dhet males (n = 14). (G) StAR protein in ovaries. Representative immunoblot of StAR, and densitometric scans of StAR immunoblots normalized to β-actin (n = 8 per group). (H) At 17 and 19 dpc, StAR immunostaining in corpus luteum of SRC-1/-2 dhet females × SRC-1/-2 dhet males (dhet) was more intense than that of WT. Three independent experiments conducted. Original magnification, ×200. Data are mean ± SEM, *P < 0.05, **P < 0.01, ***P < 0.001 (ANOVA).
LPCAT1 also catalyzes the synthesis of platelet-activating factor (PAF) from lyso-PAF and acetyl-CoA (45). PAF, an inflammatory phospholipid, was previously reported to increase in AF near term and play a role in the initiation of term and preterm labor (46–50). In the present study, we similarly observed that PAF levels in fetal lungs and AF were significantly increased at 18.5 dpc compared with earlier time points in fetuses from WT crosses (Figure 4, E and F) but failed to increase in lungs and AF of SRC-1/-2 dKO fetuses at 18.5 dpc (Figure 4, G and H). By contrast, the related acyltransferase enzyme, LPCAT2, believed to catalyze PAF synthesis in response to proinflammatory stimuli, as well as the PAF-degrading enzyme, PAF acetylhydrolase (PAFAH), were unaffected by SRC-1/-2 double deficiency in fetal lung (Supplemental Figure 5, D and E). Besides LPCAT1, expression of several enzymes associated with the de novo pathway of PAF biosynthesis (51), including ABCD2 (also known as ALDRP), a peroxisomal ATP-binding cassette transporter, which is suggested to be involved in the transport of very-long-chain acyl-CoA (52), also was analyzed; however, no differences were observed in the lungs of SRC-1/-2 dKO or dhet fetuses compared with WT fetuses (Supplemental Figure 5F).

These findings provide compelling evidence that the fetus signals the mother when its lungs develop the capacity to produce sufficient surfactant to sustain air breathing and that SRC-1 and SRC-2 serve a crucial and specific role through transcriptional regulation of SP-A and LPCAT1.

SRC-1 and SRC-2 enhance Lpcat1 promoter transactivation by glucocorticoids in a GRE-dependent manner. As mentioned above, we previously observed that SRC-1 and SRC-2 serve important roles in transcriptional upregulation of SP-A gene expression in fetal lung type II cells (20, 21). Although the Lpcat1 gene was cloned from mouse fetal lung 9 years ago (43, 44) and found to be highly expressed in lung type II cells and regulated by glucocorticoids (43), the mechanisms underlying its transcriptional regulation are largely undefined. Using the MatInspector program, we identified 5 putative glucocorticoid response elements (GREs), between −1,000 and −2,000 bp upstream of the mouse Lpcat1 transcription start site, as well as a putative Pp response element and androgen response element, between −1,000 bp and the transcription start site (Figure 5A). To functionally define the glucocorticoid-responsive region(s), different lengths of Lpcat1 5′-flanking sequence (2.0 kb, 1.0 kb, 0.6 kb, 0.4 kb) were subcloned into the pGL4.12 luciferase vector and cotransfected into lung adenocarcinoma A549 cells, together with Renilla luciferase plasmid. Cells expressing these Lpcat1 promoter constructs, as well as empty vector, were treated with or without dexamethasone (Dex, 10−7 M) for 24 hours, and luciferase activity was analyzed using a Dual Luciferase Reporter Assay System. Luciferase activities of all Lpcat1 promoter constructs were comparably increased above empty vector (Supplemental Figure 6); however, only the Lpcat1−2000-Luc plasmid containing all 4 putative GREs was upregulated by Dex treatment (Figure 5A). Moreover, when A549 cells were transfected with Lpcat1−2000-Luc reporters, with or without mutation in GRE4 and GRE5 and cultured with or without Dex (10−7 M) for 24 hours, mutagenesis of GRE4 and GRE5 prevented Dex induction of Lpcat1 promoter activity (Figure 5B). The roles of endogenous SRC-1 and SRC-2 in glucocorticoid induction of Lpcat1 were suggested by the finding that siRNA-mediated knockdown of endogenous SRC-1, SRC-2, or both SRC-1 and SRC-2 blocked the inductive effect of Dex on Lpcat1 promoter activity (Figure 5B).

To assess binding of endogenous glucocorticoid receptor (GR), SRC-1, and SRC-2 to the putative GREs, ChIP–quantitative PCR (ChIP-qPCR) was carried out in lung tissues from WT and SRC-1/-2 dKO fetal mice at 18.5 dpc using primers flanking each of these sites. As can be seen, endogenous GR, SRC-1, and SRC-2 bound specifically to GRE4 and GRE5 within the Lpcat1 promoter in fetal lung tissues of WT mice but not SRC-1/-2-deficient mice (Figure 5, C–E).

To determine whether the human LPCAT1 gene also is regulated by glucocorticoids and assess the roles of SRC-1 and SRC-2 in this regulation, we used primary cultures of HFL type II cells infected with lentivirus containing shRNA targeting SRC-1 and/ or SRC-2 and cultured them with or without Dex, in the absence or presence of the GR/PR antagonist, RU486. Dex significantly increased the mRNA and protein expression of LPCAT1 after 24 hours of treatment, and this was completely blocked by RU486 (Figure 5, F and G). In contrast, expression of LPCAT2 was not affected by Dex or by RU486 treatment (Supplemental Figure 7A). Efficient knockdown of either SRC-1 or SRC-2 (each was decreased by more than 95%, Supplemental Figure 7B) partially blocked the inductive effects of Dex on LPCAT1 expression; knockdown of both SRC-1 and SRC-2 had a more pronounced effect (Figure 5, H and I), again, suggesting possible compensatory upregulation of SRC-1 and SRC-2.

i.a. injection of SP-A and PAF at 17.5 dpc rescues the delayed-labor phenotype and downstream pathways in SRC-1/-2-deficient mice. We previously reported that i.a. injection of purified SP-A at 15.5 dpc caused preterm delivery of fetuses within 6 to 24 hours (6). In light of our findings that SP-A and PAF levels were decreased in AF surrounding SRC-1/-2–deficient fetuses, we evaluated whether administration of exogenous SP-A or PAF could rescue the delayed-labor phenotype of SRC-1/-2–deficient mice. SP-A (3 μg per sac), containing <3 fg LPS per microgram of SP-A (after polymyxin B treatment) (6), or methylcarbamyl PAF-C-16 (0.25 ng per sac) was injected i.a. into SRC-1/-2-deficient mice at 17.5 dpc; this corresponds to the approximate time that SP-A (6) and PAF (Figure 4F) naturally increase in AF. The amounts of SP-A and PAF injected were calculated to result in AF concentrations comparable to those normally found at term (~15 μg/ml [ref. 6] and 1.25 ng/ml [Figure 4F], respectively). A parallel group of SRC-1/-2–deficient mice were injected i.a. with an equivalent volume of sterile PBS, the solvent for both SP-A and methylcarbamyl PAF-C-16. Four of ten SRC-1/-2-deficient mice injected with SP-A delivered premature fetuses surrounded by intact amniotic membranes at 18.5 to 19.0 dpc, and the remaining 6 mice delivered at term on 19.5 dpc. Similarly, 3 of 10 mice injected with PAF delivered at 18.5 to 19 dpc, and the remaining 7 mice delivered at term on 19.5 dpc (Figure 6A). Only the left uterine horn was injected, while the right horn was left un.injected as an internal control. As reported
Figure 4. LPCAT1 expression and PAF synthesis and secretion are reduced in lungs of SRC-1/-2 double-deficient fetuses. (A) Lpcat1 mRNA in lungs of WT and SRC-1/-2 dhet (dhet) fetuses from 15.5 dpc to term (n = 5 mice per time point). (B) Representative immunoblot of LPCAT1 protein in lungs of WT or SRC-1/-2 dhet fetuses from 15.5 dpc to term (n = 2–3 per time point). (C) Lpcat1 mRNA in lungs of 18.5 dpc WT (n = 3), dhet (n = 14), dKO (n = 3), KO/het (n = 6), het/KO (n = 6), KO/WT (n = 3), het/WT (n = 13), and WT/het (n = 4) fetuses. (D) LPCAT1 protein in fetal lungs at 18.5 dpc. Representative immunoblot according to genotype, and densitometric scans of immunoblots of LPCAT1 normalized to β-actin for WT (n = 7), dhet (n = 8), KO/het (n = 8), and dKO (n = 4) fetuses are shown. (E) PAF in WT fetal lungs at 15.5 (n = 3), 16.5 (n = 3), 17.5 (n = 7), 18.5 (n = 5), and 19.5 (n = 3) dpc. (F) PAF in AF surrounding WT fetuses at 15.5 (n = 3), 16.5 (n = 3), 17.5 (n = 7), 18.5 (n = 6), and 19.5 (n = 4) dpc. (G) PAF in 18.5 dpc fetal lungs of WT (n = 5), SRC-1 KO (n = 4), SRC-2 KO (n = 3), dhet (n = 7), and dKO (n = 6). (H) PAF in AF surrounding 18.5 dpc WT (n = 6), SRC-1 KO (n = 4), SRC-2 KO (n = 4), dhet (n = 6), and dKO (n = 5) fetuses. (F and H) In the box-and-whisker plots, horizontal bars indicate the medians, boxes indicate 25th to 75th percentiles, and whiskers indicate 10th and 90th percentiles. Data are mean ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001, *P < 0.05, **P < 0.01 compared with WT at same dpc (ANOVA).
Figure 5. SRC-1 and SRC-2 enhance Lpcat1 promoter transactivation and expression by glucocorticoids in a GRE-dependent manner. (A) Lpcat1-luciferase reporter constructs containing various amounts of 5′-flanking sequence of the mouse Lpcat1 gene; putative GREs, androgen response elements (ARE), and P1 response element (PREs) are indicated. A549 cells cotransfected with these constructs and Renilla luciferase were treated with or without Dex (10^{-7} M) for 24 hours; luciferase assays were performed (n = 4 for each construct). (B) Effects of mutation of GRE4 and GRE5 (Lpcat1-2000MUT-Luc) and knockdown (KD) of SRC-1, SRC-2, or SRC-1 and SRC-2 on Dex-induced Lpcat1 promoter activity (n = 4, each construct). Data are mean ± SEM. (C–E) ChIP-qPCR of WT or SRC-1/-2 dKO (KO) fetal lung tissues using antibodies to (C) GR, (D) SRC-1, or (E) SRC-2 and specific primers for GRE1/2, GRE3, GRE4, or GRE5 within the Lpcat1 promoter (n = 4). (F) LPCAT1 mRNA levels in HFL type II cells treated with Dex with or without RU486 (n = 5). (G) Representative immunoblot of LPCAT1 protein in HFL type II cells treated with Dex with or without RU486. Five independent experiments were conducted. (H) LPCAT1 mRNA in HFL type II cells infected with recombinant lentiviruses expressing SRC-1 and/or SRC-2 shRNA and cultured with or without Dex (10^{-7} M) (n = 4). NTC, silencer negative control siRNA. (I) Representative immunoblot of LPCAT1 protein in HFL type II cells infected with recombinant lentiviruses expressing SRC-1 or SRC-2 shRNA, cultured with or without Dex (n = 4). Data are mean ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001; **P < 0.01 compared with Dex treatment alone, ****P < 0.001 compared with cells expressing WT Lpcat1-2000-Luc and Silencer negative control siRNA (ANOVA).
AKR1B3 (Figure 7A and B). Meanwhile, the increased expression of the PGF,α-metabolizing enzyme, PGDH, in myometrium of SRC-1/-2 dhet females bred to SRC-1/-2 dhet males also was prevented by SP-A or PAF injection (Figure 7C and D); this resulted in an upregulation of PGF2α levels in myometrium (Figure 7E). Consequently, i.a. SP-A or PAF injection caused a decrease in ovarian StAR protein (Figure 7F) and circulating P4 (Figure 7G) to levels comparable to those of WT mice. By contrast, PBS injection had no effect on any of these parameters. Again, Star mRNA expression in ovaries of SRC-1/-2 dhet females bred to SRC-1/-2 dhet males was previously for SP-A (6) and as observed in the present study for SP-A or PAF treatment, only the fetuses in the injected horn delivered prematurely — or on time in the case of SRC-1/SRC-2 double deficiency. Only 2 of 9 SRC-1/-2-deficient mice injected i.a. with sterile PBS manifested normal term labor (19.5 dpc), whereas the remaining 7 delivered 1 to 2 days late (Figure 6A).

Furthermore, we found that SP-A or PAF injection significantly reversed the inhibitory effects of SRC-1/-2 deficiency on myometrial NF-κB activation (Figure 6, B and C), contractile gene expression (Figure 6, D–G), and the expression of PGF,α synthase.
not significantly altered by SP-A or PAF injection (Supplemental Figure 3B). Based on these findings and the correlation of gestation length with the proportion of SRC-1/-2 deficient pups in the litter (Figure 1C), we postulate that secretion of both SP-A and PAF by the maturing fetal lung near term provides a key stimulus for activation of contractile genes in the maternal uterus. Supplementation with either factor was sufficient to rescue the delay in labor in SRC-1/-2 deficient mice.

Discussion

Mice in which SRC-1 and SRC-2 have both been knocked out die at birth of respiratory distress (29). To investigate the mechanisms whereby deficiency of SRC-1 and SRC-2 cause decreased production of pulmonary surfactant components, we crossed SRC-1/-2 dhet mice. Surprisingly, the mice exhibited a highly pronounced delay in parturition. Noteworthy, WT females bred to SRC-1/-2 double-deficient males (1-KO/2-het) manifested an equivalently severe delay in labor. Notably, the proportion of “double-deficient” fetuses produced by crossing WT females with 1-KO/2-het males was estimated to be similar to that of the SRC-1/-2 dhet crosses (50% vs. 56.25%, respectively), even though there are no SRC-1/-2 dKO fetuses produced in the former. Moreover, SP-A and PAF levels were significantly and comparably decreased in lungs and AF samples from all double-deficient (SRC-1/-2 dhet, 1-het/2-KO, 1-KO/2-het, and SRC-1/-2 dKO) fetuses compared with those in single-deficient (SRC-1–/– and SRC-2–/–) or WT pups. These findings strongly suggest that the delay in parturition was caused by a deficiency in fetus-derived factors in lungs of the SRC-1 SRC-2 double-deficient pups. The role of the fetus was further supported by the positive correlation found between the proportion of SRC-1/-2 deficient pups in the litters and gestational length.

Although a number of reproduction-related phenotypes have been elucidated in SRC-1 and SRC-2 KO mice (24), the combined and cooperative functions of SRC-1 and SRC-2 in pregnancy and parturition have not been explored. The present finding that double deficiency in SRC-1 and SRC-2 is required for manifestation of the delayed parturition phenotype suggests the presence of compensatory mechanisms and cooperative physiological functions between SRC-1 and SRC-2 in the regulation of gestation length. This is consistent with a previous study showing that endogenous SRC-2 overexpression in certain tissues of SRC-1–null mutants compensated partially for the loss of SRC-1 (26) as well as our current findings that KO of one coregulator in fetal lung caused marked compensatory upregulation of the other. Moreover, we observed that efficient knockdown of both SRC-1 and SRC-2 had a more pronounced effect to reduce Dex-induced LPCAT1 expression than knockdown of either SRC-1 or SRC-2 alone.

Both term and preterm labor in humans and rodents are mediated by an inflammatory response, associated with elevated levels of proinflammatory cytokines in AF and increased leukocyte migration into the myometrium (11). Increasing evidence suggests that the developing fetus may provide physical and hormonal signals that activate inflammatory transcription factors in myometrium and contribute to the induction of spontaneous labor at term. Previously, we suggested that the major surfactant protein, SP-A, a developmentally regulated C-type lectin involved in innate immunity within the lung alveolus, plays an important role in initiating labor at term by activating fetal macrophages and promoting their migration to the maternal uterus (6). Moreover, mice that were globally deficient in the genes encoding SP-A and the related C-type lectin, SP-D, manifested a significant delay in parturition, accompanied by decreased AF macrophage activation and a marked reduction in myometrial proinflammatory and contraction-associated gene expression (15).

As mentioned, SP-A transcription in fetal lung type II cells was previously found to be enhanced by SRC-1 (21) and SRC-2 (20). Accordingly, in the present study, SP-A mRNA and protein expression and secretion into AF were observed to be significantly reduced in lungs of fetuses deficient in both SRC-1 and SRC-2. The marked delay in parturition also was associated with decreased myometrial NF-κB activation and a lack of induction of the contraction-associated genes Cx43, Oxrtr, and Akr1b3 near term. Consequently, ovarian StAR protein levels failed to decline near term and circulating P levels remained elevated, due to a block in luteolysis.

While reduced SP-A expression in lungs of SRC-1/-2 deficient fetuses likely contributed to the delay in parturition, the effect of SRC-1/-2 deficiency on parturition timing (~38-hour delay) was considerably greater than that observed with SP-A or SP-A/SP-D dKO mice (~12-hour delay) (15). These pronounced differences suggested that other fetal signals contribute to the timing of labor. Previous studies by Johnston and colleagues suggested that PAF, produced in increasing amounts by the developing fetal lung and secreted into AF, may play a role in the initiation of term and preterm labor (46–50). Importantly, we observed that levels of PAF in fetal lungs and AF of SRC-1/-2 double deficient mice failed to increase and were significantly reduced as compared with those of WT or SRC-1 or SRC-2 single-deficient fetuses near term. PAF has been observed to enhance cytokine production and migration of polymorphonuclear leukocytes (PMNs) into the cervix at the time of parturition (53–55). Studies with PAF receptor KO mice suggested that PAF is a direct mediator of PMN activation (56). Moreover, unlike other inflammatory mediators (e.g., cytokines or LPS), PAF is a direct uterotonic agent (57–60). PAF also has been implicated in preterm parturition because its levels are increased in the AF of women in preterm labor (47, 61). Thus, PAF and SP-A produced by fetal lung type II cells may cooperatively act to enhance migration of macrophages and other leukocytes to the myometrium near term. Notably, we observed that i.a. injection of PAF or SP-A at 17.5 dpc corrected the parturition delay in SRC-1/-2 deficient mice. Moreover, rescue of the parturition delay in SRC-1/-2 dhet females bred to SRC-1/-2 dhet males by SP-A or PAF i.a. injection at 17.5 dpc restored NF-κB activation, Cx43 and OXTR expression in the maternal myometrium, StAR expression in the ovary, and circulating P levels to WT values at 18.5 dpc.

In studies to define the underlying mechanisms for PAF upregulation in fetal lung near term and the effects of SRC-1/-2 deficiency, we discovered that LPCAT1, a key enzyme in the acylation of sn-2 position of both DPPC and PAF (42–44), was significantly decreased in lungs of SRC-1/-2 dKO fetuses compared with that in WT fetuses. LPCAT1 was previously reported to be developmentally upregulated in mouse fetal lung, selectively expressed in type II cells, and increased by glucocorticoids (43). In studies using HFL type II cells, we observed that Dex induced LPCAT1 expression in a GR/SRC-1/-2-dependent manner. Glucocorticoid upreg-
Figure 7. Rescue of parturition delay in SRC-1/SRC-2 dhet females bred to SRC-1/-2 dhet males is associated with increased myometrial PGF$_2$α synthesis and decreased ovarian P4 production. (A) Akr1b3 mRNA (n = 6 per group) and (B) protein (n = 3 for WT, PBS, and SP-A; n = 4 for PAF) in myometrium. The same immunoblot of β-actin was used for normalization of PGDH (D), which was probed on the same blot. (C) PGDH mRNA (n = 6 per group) and (D) protein (n = 3 for WT, PBS, and SP-A; n = 4 for PAF) in myometrium. (E) PGF$_2$α levels in myometrial homogenates for WT or SRC-1/-2 mice injected i.a. with PBS, SP-A, or PAF (n = 8 per group). In the box-and-whisker plot, horizontal bars indicate the medians, boxes indicate 25th to 75th percentiles, and whiskers indicate 10th and 90th percentiles. (F) STAR protein in ovaries of WT (n = 3) or SRC-1/-2 dhet females × SRC-1/-2 dhet males injected i.a. with PBS (n = 3), SP-A (n = 3), or PAF (n = 4). (G) Maternal serum P4 levels, measured by ELISA, of 18.5 dpc WT females × WT males or SRC-1/-2 dhet females × SRC-1/-2 dhet males injected i.a. with PBS, SP-A, or PAF (n = 8 per group). (H) Proposed cooperative roles of SRC-1 and SRC-2 in the initiation of labor through regulation of SP-A and Lpcat1 gene expression. Data are the mean ± SEM. *P < 0.05, **P < 0.01, ***P < 0.001, compared with WT; *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001 compared with PBS injection (ANOVA).
ulation of LPCAT1 promoter activity was mediated by two consensus GRE repeats that bound GR, SRC-1, and SRC-2 in a cooperative manner. Previous findings that SRC-1 and SRC-2 KO mice manifested hypogonadism to glucocorticoids (62) further suggest that compromised action of glucocorticoids in SRC-1/-2-deficient mice may contribute to decreased LPCAT1 expression. Fetal plasma glucocorticoids increase exponentially during late gestation (14) and are thought to serve a role in maturation of the fetal lungs (63–68). Neonatal mice in which SRC-1 and SRC-2 were both knocked out were reported to have diffusen lung atelectasis and died at birth of apparent respiratory distress (29). Importantly, we observed that the decreased expression of LPCAT1 in lungs of SRC-1/-2-deficient fetuses was associated with profoundly reduced secretion of DPPC into AF. Since DPPC is the most surface-active component of pulmonary surfactant, this decline in DPPC production is likely responsible for the alveolar collapse observed in SRC-1/-2 dKO mice. The importance of LPCAT1 in fetal lung is supported by the finding that mice homozygous for a hypomorphic allele of Lpcat1 manifested varying perinatal mortality from respiratory distress, associated with a deficiency in surfactant DPPC (42). In light of the pleiotrophic effects of SRC-1 and SRC-2, it is likely that the actions of other transcription factors are compromised in lungs of the double-deficient mice, contributing to the impaired production of surfactant components. While our study revealed important roles of SP-A and PAF in signaling the initiation of labor, it is likely that SRC-1 and SRC-2 have additional fetal target genes besides SP-A and LPCAT1 that might contribute to the parturition delay. Genome-wide RNA-seq analysis of lung tissues of SRC-1/-2 mice will be used to identify other possible target genes and provide a more comprehensive view of the coordinated roles of SRC-1 and SRC-2 in the timing of labor.

In conclusion, our findings provide compelling evidence that the fetus initiates parturition by signaling the mother when its lungs develop the capacity to produce sufficient surfactant lipid-protein to sustain air breathing. This likely occurs through the production of two fetal lung-derived factors — SP-A and PAF (Figure 7H). Both of these factors have the capacity to activate immune cells, resulting in an enhanced inflammatory response within the maternal uterus, as well as a decline in PR function, which collectively result in increased expression of myometrial contractile genes and the initiation of labor. The coactivators SRC-1 and SRC-2 serve crucial roles in this process through transcriptional regulation of genes encoding SP-A and LPCAT1, a key regulatory enzyme in the synthesis of DPPC, which is the major surface-active component of surfactant, and PAF, a proinflammatory glycerophospholipid (Figure 7H).

Methods

Timing of labor in SRC-1/-2-deficient and WT mice. To examine the effects of deficiencies in SRC-1 and/or SRC-2 on the timing of labor, SRC-1 homozygous KO (SRC-1−/−), SRC-2 heterozygous (SRC-2−/−), and SRC-1/-2 dhet females (B6 background) were bred to genetically matched males. To determine whether phenotypic effects were caused by deficiencies in the fetuses, WT females also were bred to 1-KO/2-het males. WT B6 females were bred to WT males as controls. Breeding pairs were placed together at 1800 hours and separated the following morning at 0600 hours. Pregnancy was suggested by the presence of a vaginal plug; gestational age was designated at that time as 0.5 dpc. The time to labor was documented upon delivery of the first pup or by the presence of a litter. All mice were housed under pathogen-free conditions, maintained on a 12-hour-dark/12-hour-light cycle, and allowed free access to a standard pellet chow.

Collection of AF, myometrium, ovaries, sera, and fetal lungs at 15.5 to 18.5 dpc. Parturition timing was assessed in first pregnancies, while fluids and tissues were analyzed in second pregnancies. AF, myometrium, ovaries, sera, and fetal lungs were collected from timed-pregnant mice at approximately 1000 hours at 15.5, 16.5, 17.5, and 18.5 dpc. Uteri were exposed and AF from individual amniotic sacs was carefully aspirated, avoiding maternal blood contamination, using a 20-gauge needle and a 1.0-ml syringe. Maternal myometrium was isolated after removal of all fetus-derived tissues, followed by gentle scraping and blotting to remove the endometrial layer. Ovaries were dissociated from surrounding fat. Intact fetal lungs were harvested on ice. AF from each sac and lungs from each fetus were separately harvested for subsequent assignment of genotype. Maternal blood was collected at 18.5 dpc by retro-orbital puncture of anesthetized animals and allowed to clot for 2 hours at room temperature before centrifuging for 20 minutes at 3,000 g. Sera were harvested and stored at –80°C. All tissues were rinsed in ice-cold 1× PBS, flash frozen in liquid nitrogen, and stored at –80°C until analysis. Forelimbs from each fetus were placed in phenol-chloroform for DNA isolation and genotyping.

Animal surgery for injection of substances into AF. Timed-pregnant mice at 17.5 dpc were anesthetized with isoflurane via a precision vaporizer. The left uterine horn was gently pulled through a 1-cm incision made above the visible ovarian fat pad; SP-A (3.0 ng in 50 μl PBS), methylcarbamyl PAF-C-16 (Sigma-Aldrich, 0.25 ng in 50 μl PBS), or PBS (50 μl) was injected through the exposed uterine wall into each amniotic sac with a sterile 31-gauge half-inch needle (Becton Dickinson). The right uterine horn was untouched. The left uterine horn was returned to the abdominal cavity, the abdominal muscle wall was closed with Suturevet 4/0 polyglycolic acid synthetic absorbable sutures (Butler Schein Animal Health), and the skin was closed by using 7.5-mm wound clips (Perfect Agrafes Chirurgicales). The mice were kept on a warming pad and returned to their cages after recovery. One day later (18.5 dpc), mice were sacrificed, and myometrium, ovaries, and sera were harvested, as described above, or the mice were allowed to deliver for analysis of parturition timing.

Cell culture. Midgestation HFL tissues were obtained from Advanced Bioscience Resources. HFL type II pneumocytes were isolated and cultured as described in detail previously (69). Briefly, the fetal lung tissues were minced and rinsed in serum-free Waymouth’s MB752/1 medium (Invitrogen). Lung explants were plated on lens paper supported by stainless steel grids in 100-mm sterile dishes containing 3 ml serum-free Waymouth’s medium containing Bt cAMP (1 mM) (Roche) for 3 to 5 days to enrich the population of differentiated type II cells. Cells were dispersed from the explants by digestion with collagenases type I (0.5 mg/ml; Sigma-Aldrich) and type IA (0.5 mg/ml; Sigma-Aldrich) for approximately 10 minutes. The resulting cell suspension was depleted of fibroblasts by incubation with diethylaminoethyldextran (250 μg/ml) for 30 minutes at 37°C, followed by centrifugation at 400 g for 5 minutes. The cell pellet was resuspended in Waymouth’s MB752/1 medium containing 10% (vol/vol) FBS (Gemini Bio-Products), plated onto 6-well culture plates (1.5 × 10⁶ cells per well) coated with extracellular matrix prepared from Madin-Darby
canine kidney cells (CRL 6253; ATCC) (69) and incubated overnight. Cultured cells were washed twice with medium to eliminate dead and nonadherent cells. Cells were then infected with lentiviruses containing shRNAs targeting SRC-1 and/or SRC-2 overnight and treated with or without Dex (10^{-7} M) in phenol red-free DMEM medium without FBS, with or without RU486. The human lung adenocarcinoma cell line A549 (CCL 185, ATCC) was maintained in Waymouth’s MB752/1 medium containing FBS (10% vol/vol).

**Lentivirus generation and infection.** Lentiviral vectors containing shRNAs targeting SRC-1 or SRC-2 in pGIPZ vectors were obtained from a lentivirus shRNA library (Open Biosystems). Two different shRNA vectors were used for silencing SRC-1 (NCOA1, catalog RHS4430-99148014) and SRC-2 (NCOA2, catalog RHS4430-9914731), and their efficacy was validated. Lentiviral supernatants were produced in human embryonic kidney 293T (HEK 293T) cells by Polyethylenimine-mediated (Polysciences) transfection of plasmids pMD2.G, ps-PAX2, and human pGIPZ lentiviral shRNA for SRC-1 and SRC-2. Lentiviral supernatants were collected after 48 and 72 hours of culture. Viral titers were determined by measuring the coexpressed GFP after infection of HEK 293T cells.

HFL type II cells were then infected with recombinant lentiviruses containing shRNAs targeting SRC-1 and/or SRC-2 overnight and treated with Dex (10^{-7} M) in phenol red-free DMEM medium without FBS, in the absence or presence of RU486. For control experiments, cells were infected with nonsilencing pGIPZ lentiviral shRNA.

**SP-A isolation.** Human alveolar proteinosis lavage was delipidated using isopropyl ether and 1-butanol (6). Serum albumin was removed by extraction on a DEAE-Affigel blue gel column (Bio-Rad). Endotoxin content was assessed using a Limulus Amebocyte Lysate endotoxin assay kit (Cambrex). Remaining endotoxin content was measured using an antibody raised against SP-A (70).

**Quantitative RT-PCR.** Total RNA from tissues or cells was extracted using an RNaseasy Mini Kit (Qiagen). The purity and integrity of the RNA were checked spectrophotometrically and by gel electrophoresis before use. Quantification of total RNA was performed by measuring absorbance at OD_{260}. RNA was treated with DNase (Invitrogen) to remove any contaminating DNA, and 2 μg were reversed transcribed using the SuperScript III cDNA First-Strand Synthesis Kit (Invitrogen). Primer sets are listed in Supplemental Table 3. All primers were validated for efficiency by creating standard curves using a gradient of diluted samples with r² values larger than 0.96. The specificity of the primers was verified by examining the melting curve. Product purity was confirmed by gel electrophoresis. For quantitative analysis of mRNA, a CFX384 Real-Time PCR Detection System (Bio-Rad) was employed using Power SYBR Green PCR Master Mix (Applied Biosystems) for the detection of PCR products. The cycling conditions were 50°C for 2 minutes and 95°C for 10 minutes, followed by 45 cycles of 95°C for 15 seconds, and 60°C for 1 minute. As a negative control for all of the reactions, distilled water was used in place of cDNA. Amplification of the housekeeping gene 36B4 was determined for sample loading and normalization. Each sample was amplified in triplicate, and the mean of Cq value was calculated. The relative fold changes were calculated using the comparative Ct method (2^ΔΔCt) (71).

**Immunoblot analysis.** Total protein extracts were prepared from tissues using RIPA buffer (Cell Signaling). Nuclear and cytoplasmic proteins were separated by NE-PER Nuclear and Cytoplasmic Extraction Reagents (Thermo Scientific). Equivalent amounts of protein determined by BCA assay (Thermo Scientific) were resolved by NuPAGE 4-12% Bis-Tris gel (Invitrogen) electrophoresis and blotted to Hybond-P PVDF membranes (Amersham Biosciences). The following primary antibodies and dilutions were used: anti–SP-A (1:500; ref. 70); anti–LPCAT1 (1:3,000; ab94903, Abcam); anti–SRC-1 (1:1,000; A300-344A, Bethyl Laboratories); anti–SRC-2 (1:1,000; A300-346A, Bethyl Laboratories); anti–NF-κB p65 (1:500; SC-372, Santa Cruz Biotechnology); anti–NF-κB p50 (1:500; SC-114, Santa Cruz Biotechnology); anti–OXTR (1:500; SC-33209, Santa Cruz Biotechnology); anti–AKR1B3 (1:500; SC-271007, Santa Cruz Biotechnology); anti–PGDH (1:500; SC-98907, Santa Cruz Biotechnology); anti–COX-2 (1:1,000; ab15191, Abcam); anti–20α-HSD (1:5,000; gift of Geula Gibori, University of Illinois, Chicago, Illinois, USA); anti–StAR (1:300; ab58013, Abcam); anti–β-actin (1:5,000; ab8227, Abcam); anti–GAPDH (1:2,000; ab9485, Abcam); and anti–histone H3 (1:2,000; 05-928, Millipore). Horseradish peroxidase–conjugated anti-rabbit or anti-mouse IgG (GE Healthcare) was used as secondary antibody. The membranes were developed using Supersignal West Pico Chemiluminescent substrate (Thermo Scientific). To control for sample loading and transfer, the ratio of band intensities to β-actin (for total protein; ab8227, Abcam), GAPDH (for cytoplasmic protein; ab9485, Abcam), or histone H3 (for nuclear protein; clone A35 05-928, Millipore) was obtained to quantify the relative protein expression level.

**Plasmid and siRNA transfection and luciferase assays.** Various amounts of DNA flanking the transcription start site of the mouse Lpcat1 gene were amplified from mouse genomic DNA and subcloned into the pGL4.12 vector (Promega) to generate luciferase reporter plasmids. Site-directed mutagenesis of putative GREs was performed using the QuickChange II Site-Directed Mutagenesis Kit (Agilent Technologies). All constructs and mutations were confirmed by DNA sequencing at the McDermott Center Sequencing Core, University of Texas Southwestern Medical Center. SRC-1 and SRC-2 transcripts in A549 cells were knocked-down by transfection of preannealed Silencer Select siRNA duplexes (s16461 for SRC-1 and/or s20528 for SRC-2, Ambion) using Lipofectamine RNAiMAX transfection reagent (Invitrogen). Silencer Negative Control #1 siRNA (AM4611, Ambion) was transfected as a negative control. The efficiency of knockdown was determined by RT-PCR and Western blotting. Twenty-four hours after transfection, WT, SRC-1, SRC-2, or SRC-1/-2 double-knockdown A549 cells were transfected with various Lpcat1-luciferase reporter constructs and Renilla luciferase plasmid using Lipofectamine 2000 transfection reagent (Invitrogen) according to the manufacturer’s protocol. Dex (10^{-6} M) was added into the medium for 24 hours and then whole-cell lysates were prepared using Passive Lysis Buffer (Promega). Luciferase assays were performed using the Dual Luciferase Reporter Assay System (Promega), according to the manufacturer’s protocol, in a GloMax plate-reading luminometer (Promega).

**ChIP-qPCR.** ChIP assays were performed on minced mouse fetal lung using a ChIP Assay Kit (Millipore) as described previously (72). Cross-linking was performed using formaldehyde (1% in PBS) at room temperature for 10 minutes with gentle rotation. The minced tissues were pelleted in a microfuge at full speed for 1 minute at 4°C. The pellets were washed, resuspended in SDS lysis buffer containing protease inhibitor cocktail, and then sonicated. Sheared chromatin was diluted in ChIP...


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The role of steroid receptor coactivator 2 (SRC-2) in the regulation of mouse reproduction is distinct from the role of SRC-1. SRC-2 is essential for the proper function of the steroid receptor coactivator (SRC) family.


