CC chemokine ligand 2 (CCL2) stimulates aromatase gene expression in mammary adipose tissue

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Abstract

Obesity is a risk factor for postmenopausal breast cancer. Obesity-related inflammation upregulates aromatase expression, the rate-limiting enzyme for estrogen synthesis, in breast adipose tissue (BAT), increasing estrogen production and cancer risk. The regulation of aromatase gene (CYP19A1) in BAT is complex, and the mechanisms linking obesity and aromatase dysregulation are not fully understood. An obesity-associated factor that could regulate aromatase is the CC chemokine ligand (CCL) 2, a pro-inflammatory factor that also activates signaling pathways implicated in CYP19A1 transcription. By using human primary breast adipose stromal cells (ASCs) and aromatase reporter (hARO-Luc) mouse mammary adipose explants, we demonstrated that CCL2 enhances the glucocorticoid-mediated CYP19A1 transcription. The potential mechanism involves the activation of PL4 via ERK1/2 pathway. We also showed that CCL2 contributes to the pro-inflammatory milieu and aromatase expression in obesity, evidenced by increased expression of CCL2 and CYP19A1 in mammary tissues from obese hARO-Luc mice, and subcutaneous adipose tissue from obese women. In summary, our results indicate that postmenopausal obesity...
1 | INTRODUCTION

Adult obesity is a well-known risk factor for developing the most commonly diagnosed cancer among women, estrogen receptor positive (ER+) postmenopausal breast cancer. After menopause, estrogens are produced from androgens by the enzyme aromatase in peripheral tissues, adipose tissue being the major source. Excessive fat accumulation associates with a chronic low-grade inflammation, and elevated aromatase expression and activity in breast adipose tissue (BAT) of post-menopausal women. Combined, these changes are seen as the major driving forces increasing local estrogen production and promoting breast carcinogenesis after menopause.

Human aromatase (CYP19A1) gene expression is controlled by alternative usage of at least nine tissue-specific promoters, each of which is expressed and regulated in a tissue-selective manner by distinct set of hormones, cytokines, and/or growth factors. In the healthy BAT, basal levels of CYP19A1 mRNA transcripts are thought to be maintained by the exclusive use of the aromatase promoter I.4 (PI.4) and the balance between the transcriptional inducers and repressors of this promoter within the tissue. Upregulation of PI.4 in adipose tissue is closely linked with the local pro-inflammatory factors, tumor-necrosis factor (TNF)-α, interleukin (IL)-6, IL-1β, and IL-8, produced by hypertrophic adipocytes and resident macrophages. Glucocorticoids, always present in BAT, are essential for these factors to induce PI.4 and CYP19A1 expression in adipose stromal cells. On the contrary, anti-inflammatory factors, such as IL-10 produced by, for example, mammary parenchyma and M2 macrophages, mediate its transcriptional repression. In postmenopause, BAT inflammation is a common phenomenon, indicated by increased number of macrophages and crown-like structures (CLS, necrotic adipocytes surrounded by macrophages). Although this process is primarily driven by postmenopausal weight gain, it may also occur in women with normal BMI. CCL2, also known as monocyte chemotactic protein-1 (MCP-1), is a key factor in macrophage-mediated inflammation in the obese white adipose tissue. It is widely recognized by its potent chemoattractant effect when binding to its receptor on monocytes, macrophages, and lymphocytes. High circulating and tissue levels of CCL2 have been found in obese subjects, as well as in women with breast cancer. and its receptor CCR2, a G-protein coupled receptor, are expressed by a large variety of cells in subcutaneous fat including mature adipocytes, fibroblasts, and immune cells, such as macrophages. Interestingly, CCL2-CCR2 interaction involves the activation of the NF-κB and MAPK-ERK1/2 cellular pathways, both of them implicated in the transcription of aromatase PI.4 in BAT. Given the importance of CCL2 in supporting the inflammatory environment and its potential in activating molecular pathways that regulate aromatase PI.4, we hypothesize that CCL2 could also play a direct role in modulating the local expression of aromatase in BAT. In this study, we focused on investigating the effects of adiposity and CCL2 on aromatase gene expression in the mammary tissue by using a transgenic human aromatase reporter mouse (hARO-Luc mouse) model, and subcutaneous adipose tissue samples and breast adipose stromal cells from women.

2 | MATERIALS AND METHODS

2.1 | Animals

Animal care and use was conducted in accordance with the Finnish Act on the Use of Animals for Experimental Purposes and EU laws, guidelines, and recommendations. The study protocols and procedures were approved by the national Animal Experiment Board in Finland (ESAVI/7471/04.10.03/2012).

The hARO-Luc mice expressing the full regulatory region of the human aromatase gene (CYP19A1) were housed under standard conditions in the Central Animal Laboratory at the University of Turku. The mice were maintained with 12 hours light/dark cycle in constant temperature (21 ± 3°C) and humidity (55 ± 15%), fed with soy-free RM3 chow (SDS, Whitham, Essex, UK) and tap water ad libitum. Mouse line maintenance and genotyping was done in collaboration with the Turku Center for Disease Modeling (TCDM).

Six-week-old hARO-Luc female mice were allocated into two groups with similar body weights and littermate distribution, and fed with purified low-fat D12450B diet (LFD, 10% of calories from fat) or high-fat D12492 diet (HFD, 60% of calories from fat) obtained from Research Diets Inc (New Brunswick, NJ). Experiments were done in three parts with 4 to 6 animals per group for a final number...
of 16 mice on LFD and 17 mice on HFD. After 8 weeks, whole body fat content was measured with EchoMRI (Echo Medical Systems, Houston, TX, USA) and the mice were sacrificed with CO₂ followed by neck dislocation. Four hours fasting tail vein blood was collected for immediate fasting glucose analysis and heart puncture blood for serum. Inguinoabdominal mammary gland (lymph node removed) samples were collected to liquid nitrogen for the aromatase reporter (luciferase) activity measurements. Different adipose tissue depot samples (mammary gland, gonadal fat, and retroperitoneal fat) were collected to culture medium for ex vivo tissue-derived cytokine measurements (described below), and also to 10% of neutral-buffered formalin for histological analyses.

### 2.2 | Ex vivo mammary adipose tissue cultures

As previously described by McGillicuddy et al., inguinoabdominal mammary gland, gonadal and retroperitoneal fat samples from hARO-Luc mice were divided into explants of approximately 100 mg each, and placed separately on 24-well culture plates containing 1 mL of Modified Eagle Medium (DMEM/F12, Life Technologies Ltd, Grand Island, NY, USA) supplemented with 100 IU/mL of penicillin and 100 μg/mL of streptomycin. In each experiment, 4 to 6 animals were used.

For tissue producing cytokine measurements, tissue culture media were collected after 24 hours incubation at 37°C in a humidified atmosphere with 5% CO₂, and then frozen at −70°C until analyzed. CCL2, leptin, IL-6, and TNFα concentrations were analyzed from tissue exposed media and serum samples with a MILLIPLEX MAP Mouse Adipocyte Magnetic Panel (#MADCYMAG-72K; Merck Millipore, Billerica, MA, USA) according to manufacturer's instructions by using the Luminex 200 analyzer (Luminex Corporation, Austin, TX, USA).

To investigate the potential modulation of luciferase (Luc) aromatase reporter activity by CCL2 in mammary gland explants, the antibiotics containing serum-free DMEM/F12 medium was supplemented with 250 nM dexamethasone (DEX, SERVA Electrophoresis GmbH, Heidelberg, Germany) alone or in combination with 100 ng/mL of CCL2 (PeproTech, Rocky Hill, NJ, USA). After 24 hours incubation, mammary gland tissues were collected and analyzed for Luc activity.

### 2.3 | Luciferase reporter activity assay

The Luc reporter activity in mammary gland was measured as described before. Shortly, tissue samples were homogenized in buffer containing 25 mM of Tris acetate (pH 7.8), 1.5 of mM EDTA, 10% of glycerol, 1% of Triton X-100, 2 mM of dithiothreitol, and plus 1 x Roche Complete MiniProtease inhibitor tablets (Roche Diagnostics, Penzberg, Germany). The homogenates were then centrifuged at 800 g for 30 minutes at +4°C. Luc activity was measured from the supernatant with the Luciferase assay kit (BioThema AB, Handen, Sweden) according to manufacturer's instructions by using the Victor2 Multilabel counter (PerkinElmer, Turku, Finland). The results were related to the sample weight or protein content measured by using Pierce BCA Protein Assay Kit (Thermo Fisher Scientific, Rockford, IL, USA) according to manufacturer's instructions.

### 2.4 | Subcutaneous adipose tissue

Snap-frozen human abdominal subcutaneous adipose tissue samples were purchased from ZenBIO Inc (Research Triangle Park, NC, USA). Tissue samples were collected from nonobese BMI (≤25) or obese (≥30) women aged 32-47 years.

### 2.5 | Primary human breast adipose stromal cells (ASCs)

Primary breast ASCs isolated from breast tissues of women undergoing elective breast reduction or mastectomy were purchased from ZenBIO Inc or were obtained from Turku University Hospital (Ethical committee approval ETKM 23/2018). The patients gave their written informed consent for sample usage. ASCs were isolated from dissected human tissue by enzymatic digestion and differential pulse centrifugations as described previously. At least three independent experiments using 3 different donors cells each run in triplicates (aged 20-66 y, BMI: 25.1-29.1 kg/m²). For expansion, ASCs were plated at a density of 2x10⁶/cm² in DMEM/F12 supplemented with 5%-15% heat inactivated fetal bovine serum, 100 IU/mL penicillin and 100 μg/mL streptomycin and were maintained at 37°C humidified atmosphere with 5% CO₂. Cells were passaged once by treating with 0.25% of trypsin-EDTA and seeded (40 000 cells/well) on 12-well plates. At 70% confluency, cells were serum starved for 12 hours in medium containing 0.1% of BSA. Cells were treated with 10 nM DEX alone and in combination with 100 ng/mL CCL2 and with one of the inhibitors (10 μM): U0126 (MEK-ERK1/2 inhibitor, Sigma-Aldrich), BAY 11-7082 (NF-κB inhibitor, Sigma-Aldrich), or RS 504393 (CCR2 inhibitor, Tocris Bioscience, Bristol, UK). The effectiveness and relevance of these inhibitors to inhibit each of their target proteins have been widely demonstrated in different biological systems.
2.6 RNA isolation, cDNA synthesis, and quantitative-PCR

Total RNA was isolated from human ASCs or snap-frozen adipose tissue samples by using TRIzol (Bioline, Luckenwalde, Germany) according to the manufacturer’s instructions. One μg of total RNA was treated with deoxyribonuclease I (DNase I amplification grade, Sigma-Aldrich, St Louis, MO, USA) and converted to cDNA by using SensiFAST cDNA synthesis kit (Bioline, London, UK). Quantitative-PCR (qPCR) was performed using the Dynamo Flash SYBR Green qPCR Kit (Thermo Fisher Scientific) on the CFX96 Real-Time C1000 Thermal Cycler (Bio-Rad, Hercules, CA, USA). Primers used were: CYP19A1 F: 5′- TTGGAAATGCTGACCCGAT- 3′, CYP19A1 R: 5′- CAGGAATCTGCCGTGGGAGA- 3′, aromatase Pl.4 F: 5′- GTAGAACGTGACCAACTGG- 3′, aromatase Pl.4 R: 5′- CACCCGTTGTGATGTTGCAAGCCTGCC- 3′, CCL2 F: 5′- AATGGTCTTGAAGATCACAGCTTC- 3′ and CCL2 R: 5′- TAGCAGCCACCTTCATTCCCCAACAG- 3′. Cycling conditions were 95°C for 10 minutes and variable cycles of 95°C for 10 seconds, 59°C for 30 seconds and 72°C for 15 seconds. All samples and standards were run in triplicate. Expression levels of the final products were normalized to human housekeeping genes: ACTB F: 5′- TGCGTGACATTAAGGAGAAG- 3′, ACTB R: 5′- GCTCGTAGCTCTTCTCCA- 3′ and GAPDH F: 5′- TGGTATCGTGGAAGGACTCATGAC- 3′, GAPDH R: 5′- ATGCCAGTGAGCTTCCCGTTCAGC- 3′.

2.7 ERK1/2-MAPK activity

ASCs were plated in 24-well plates at the density of 20 000 cells/well. At 80%-90% confluency, cells were treated with one the inhibitors (10 μM): R504393 or U0126. Fifteen min later, 100 ng/mL of CCL2 was added, and after another 15 minutes, endogenous levels of phosphorylated ERK1/2 protein was measured from the cell lysates (containing 10-15 μg of protein per well) by using ERK1/2 (pT202/Y204) SimpleStep ELISA Kit (Abcam). The assay was performed according to the manufacturer’s instructions and analyzed at 450 nm using VICTOR Multilabel Plate Reader (Perkin Elmer, Turku, Finland).

2.8 Statistical analyses

Statistical analyses were performed using GraphPad Prism version 8.0.1 for windows (GraphPad Software Inc, San Diego, CA, USA). To assess the statistical significance between two groups, unpaired parametric t test was used. The differences between multiple treatments groups were tested by one-way analysis of variance (ANOVA) followed by Tukey’s or Dunnett’s multiple comparisons test. Pearson correlation test was used to evaluate the relationship between two variables. Differences were considered statistically significant at \( P \leq .05 \). Data are expressed as mean ± standard error of mean.

3 RESULTS

3.1 HFD induces weight gain and adiposity and impairs the metabolic status of female hARO-Luc reporter mice

Female hARO-Luc mice fed with HFD gained more weight and accumulated more body fat during the 8-week intervention, compared to the LFD-fed mice (Figure 1A,B). At the end of the experiment, HFD-fed mice were at least 38% heavier and had accumulated 35% more fat than mice fed with LFD and exhibited higher circulating leptin and glucose levels (Figure 1C,D).

**FIGURE 1** HFD induces obesity and obesity-related metabolic disturbances in female hARO-Luc mice. A, Body weight gain. B, Body adipose mass. C, Serum leptin levels. D, Circulating blood glucose levels. LFD N = 16 mice and HFD N = 17 mice. Data are shown as mean ± SEM and was analyzed by unpaired parametric t test. *\( P < .05 \), **\( P < .001 \)
3.2 | Weight gain and adiposity are drivers of both the increased aromatase Luc reporter activity and CCL2 levels in mammary gland of HFD-fed hARO-Luc mice

As has been shown before, HFD induces a variable weight gain in FVB/N mice.36,37 In our study, the majority of animals on HFD exhibited evident metabolic changes, but three out of 17 mice did not gain weight or, gained less than 20%, after the 8 weeks of dietary intervention. In animals with ≥20% weight gain the expression of aromatase Luc reporter in mammary fat tissue was significantly higher than in the mammary of mice that remained lean (Figure 2A). Furthermore, mammary gland hARO-Luc reporter expression correlated positively with both weight gain and adiposity (Figure 2B,C).

Interestingly, mammary Luc reporter expression correlated positively with tissue pro-inflammatory factors, CCL2 (Figure 2D), IL-6 ($R^2 = 0.72$, $P = .05$), and leptin ($R^2 = 0.60$, $P = .07$). However, only a low number (≤2 per mm$^2$) of CLS was detected in mammary fat tissue, despite the weight gain and increased adiposity. Additionally, CLS numbers did not correlate with the tissue Luc reporter expression (data not shown). Notably, in the gonadal or retroperitoneal fat tissue no significant correlation was found between CCL2 or IL-6 levels and Luc reporter activity (Supplementary Table 1). The levels of TNFα, IL-1β, IL-1α, adiponectin, VEGF, PAI-1, and resistin did not correlate with aromatase reporter expression in mammary, gonadal, or retroperitoneal fat tissue (Supplementary Table 1).

3.3 | CCL2 and CYP19A1 mRNA expressions are upregulated in subcutaneous adipose tissue of obese women

In order to confirm the findings from the mouse model, we investigated whether adipose tissue from obese and lean women show any difference in the expression of CCL2 or total CYP19A1 mRNA. Our results show that subcutaneous adipose tissue of obese women exhibited significantly higher aromatase gene expression and almost sixfold increase in CCL2 gene expression compared to those tissue samples collected from nonobese women (Figure 3A,B).

3.4 | CCL2 upregulates CYP19A1 in the breast adipose tissue through promoter I.4

To examine whether CCL2 directly modulates glucocorticoid-dependent aromatase gene expression through promoter I.4

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**FIGURE 2** In HFD-fed hARO-Luc female mice weight gain and adiposity associate with increased both Luc aromatase reporter activity and CCL2 expression in mammary gland. A, hARO-Luc reporter activity. Data are shown as mean ± SEM and was analyzed by two-way ANOVA with Tukey’s multiple comparisons post hoc test. *$P < .05$, no significant (n.s). Correlations between: B, Luc reporter activity in mammary gland and body weight gain, C, Luc reporter activity in mammary gland and body adiposity, and D, Luc reporter activity and CCL2 levels in mammary gland of HFD-fed mice. Open circles in figures B-D denote data from the same six animals. Counts per second (CPS)
in mammary adipose tissue, ex vivo cultures of hARO-Luc tissue were performed. Indeed, in the presence of DEX, addition of CCL2 caused a further stimulation of Luc reporter activity in mammary tissue explants from hARO-Luc mice (Figure 4).

Next, we investigated whether CCL2 could also regulate aromatase expression in human breast adipose tissue. In cultured primary human breast ASCs, similar to hARO-Luc mammary gland tissue, CCL2 enhanced total CYP19A1 gene expression levels (Figure 5A). CCL2-induced aromatase CYP19A1 gene transcription was prevented by blocking the CCL2 receptor CCR2, with the specific receptor antagonist RS504303 (Figure 5A). Additionally, to examine whether CCL2 mediates aromatase expression via aromatase promoter I.4, transcript levels of this promoter were quantified in ASCs after treatments. Correspondingly, CCL2 enhanced over threefold the effect of glucocorticoids on aromatase promoter I.4 transcripts expression (Figure 5B), thus indicating a specific action through this distal promoter. No significant induction of aromatase promoter II-driven aromatase expression was detected (Supplementary Figure 1).

3.5 | Activation of the MAPK-ERK1/2 signaling pathway is necessary for CCL2 effect on CYP19A1 gene expression in ASCs

The intracellular signaling pathway displayed upon CCL2-CCR2 activation on the cell surface involves both the NF-κB and MAPK inflammatory cascades. Besides of being important for modulating cell immune responses, the two pathways are also known to regulate aromatase promoter I.4 transcriptional activity in adipose stromal cells. To determine the potential implication of the two signaling pathways in mediating CCL2 effect on aromatase gene expression, specific inhibitors of both cascade proteins were tested in ASC cultures in the presence of DEX and CCL2. Our results demonstrate that inhibition of the ERK1/2 pathway with the inhibitor U0126 led to a significant reduction in the CCL2 stimulatory effect on both PI.4 and CYP19A1 expression (Figure 5B,C, respectively), whereas inhibition of the NF-κB pathway did not show significant effect on total CYP19A1 expression (Figure 5C).

The activation of the MAPK-ERK1/2 signaling pathway in primary breast ASCs by CCL2 was further confirmed using a Phospho-ERK1/2 ELISA assay. Phosphorylated ERK1/2 protein levels were measured after treatment with CCL2 alone.
or in combination with specific CCR2 inhibitor RS 504393 or with MEK-ERK1/2 inhibitor U0126. As expected, CCL2 stimulated phosphorylation of ERK1/2 intracellular pathway in breast preadipocytes (Figure 5D), while preventive treatment with CCR2 and MEK-ERK1/2 inhibitors reduced CCL2-mediated phosphorylation of ERK1/2 (Figure 5D).

4 | DISCUSSION

Inflammation and increased expression of aromatase are important risk factors for postmenopausal breast cancer. However, the regulation of aromatase (CYP19A1) gene expression in breast tissue is complex, and the mechanisms linking adipose tissue inflammation to the increase in aromatase expression are not fully understood. In this study, we demonstrated that CCL2, a chemokine secreted by adipocytes and macrophages, besides being overexpressed in WAT of obese women, also directly regulates aromatase gene expression in BAT via promoter I.4.

In response to adipose tissue expansion during obesity, hypertrophic adipocytes secrete chemokines, particularly CCL2, to recruit monocytes from the circulation, which facilitates tissue remodeling and adaptation to the altered metabolism. Increased adiposity often associates with abnormally high secretion of CCL2, which in turn, promotes monocyte recruitment and development of CLS, and augments production of pro-inflammatory factors, including TNFα, IL-6, PGE2, and CCL2, and ultimately, increases aromatase expression in BAT.39,40 Although the most important actions of CCL2 are related to chemotaxis, this cytokine also triggers several intracellular signals implicated in inflammation, angiogenesis, and carcinogenesis.41-44 Most of these intracellular signals associate with the activation of the PI3K/Akt/NF-κB cascade and the mitogen-activated protein kinase (MAPK) pathway, particularly the Ras/MEK/ERK
Interestingly, these signaling pathways have also been demonstrated to regulate aromatase promoter I.4 transcription expression in breast ASCs. Given the importance of CCL2 in macrophage-mediated BAT inflammation, as well as its potential to activate NF-κB and MAPK pathways, it is pertinent to propose that CCL2 may also regulate aromatase expression in a transcriptional level. In our study, we showed that, indeed, CCL2 activates aromatase gene transcription in the breast tissue. By utilizing hARO-Luc-derived mammary tissue explants and primary breast ASCs, we have found that CCL2 specifically induces glucocorticoid-dependent CYP19A1 expression via the activation of the distal promoter I.4. Such activation seems to be particularly mediated through its membrane receptor, CCR2, and by the following stimulation of MAPK/ERK1/2 signaling pathway in primary breast ASCs. However, we cannot exclude the possibility that other pro-inflammatory pathways in addition to MAPK/ERK1/2 and NF-κB, may also be implicated in this stimulatory effect. We then propose that CCL2 may regulate CYP19A1 transcription in similar manner than TNFα, a well-known inducer of aromatase promoter I.4 in adipose tissue. Prior studies have reported that TNFα acts cooperatively with glucocorticoids to induce CYP19A1 expression in breast ASCs, acting through NF-κB and MAPK signaling cascades, downstream stimulation of EGR2 and EGR3 transcription factors, and activation of the AP-1 element within PI.4 transcriptional region. Nevertheless, further studies are needed to determine whether transcriptional mechanisms of CCL2 involve also the activation of AP-1 responsive element.

Expression of aromatase in a noncancerous BAT either before or after menopause is primarily driven by the use of the I.4 promoter located approximately 73 kb upstream of the common coding region. In noncancerous conditions, low levels of aromatase transcripts are thought to be maintained via PI.4 thanks to normal tissue levels of pro-inflammatory factors, providing low levels of estrogens in the breast tissue. As previously established, elevated BMI in postmenopausal women positively associates with higher circulating and adipose tissue levels of estrogens, attributed, in part, to increased inflammation and aromatase expression in WAT. Adiposity and subsequent adipose tissue inflammation are known to promote a gradual increase in promoter I.4 transcripts. It has also been reported that glucocorticoid levels are higher in the obese subcutaneous adipose tissue. The increase in BAT aromatase expression in obese women is likely due to pro-inflammatory factors rather than glucocorticoids as there is no evidence of a direct association between systemic glucocorticoid and aromatase expression levels.

The magnitude of aromatase gene upregulation in BAT, however, depends, for example, on the degree of adiposity, the levels of pro-inflammatory factors and the abundance of CLS. In line with this, we observed a positive correlation between mammary aromatase reporter activity and weight gain in HFD-fed female hARO-Luc (FVB/N) mice. Interestingly, no weight-related changes in aromatase reporter expression were observed in gonadal, visceral or retroperitoneal fat, indicating that obesity-related aberration in CYP19A1 regulation is fat depot specific. This is supported by earlier studies with Arom knockout FVB/N mouse models, showing upregulation of aromatase by HFD feeding in mammary fat pad, but not in intra-abdominal fat depots. In these studies, unlike ours, higher activity of promoters I.4 and II was associated with a significant increase in CLS density, indicating a rather severe inflammatory condition. Interestingly, in our current study, similar to our earlier studies in hARO-Luc males and ovariectomized females, weight gain and adiposity-induced mammary gland aromatase reporter expression via promoter I.4 independent of CLS. This indicates that the pro-inflammatory milieu, prior the onset of CLS, may be sufficient to activate promoter I.4 and upregulate aromatase expression in breast tissue.

The inflammatory milieu and the regulation of aromatase gene transcription in subcutaneous adipose tissue differ from visceral fat depots. Despite the significant increase in fat content in visceral and subcutaneous adipose tissue in our in vivo experiments, aromatase expression and pro-inflammatory factors were significantly altered only in the mammary fat pad. And unlike visceral fat depots, levels of pro-inflammatory cytokines IL-6 and CCL2, and leptin, were significantly higher in mammary fat pad of animals with significant weight gain and correlated positively with mammary aromatase reporter activity. It is important to note that the use of ovary intact mice in this study may have attenuated the inflammation and, in turn, the levels of cytokines and aromatase in the mammary fat pad and other fat tissue depots. In obese women, level of CCL2 is increased in WAT, where it also associates with macrophage infiltration and insulin resistance. Furthermore, circulating CCL2 increases in postmenopause, independent of BMI, as well as in serum and tumors of women with breast cancer. In this study, the expression levels of CCL2 and aromatase transcripts in subcutaneous adipose tissue of obese women were higher, compared to the adipose tissue of lean women, supporting the finding of CCL2 as a factor that induces adipose aromatase expression.

High circulating level of estrogens is accepted as a marker of increased risk for developing postmenopausal breast cancer. Estrogen concentrations in the bloodstream after menopause reflect the metabolism and production rates of estrogens in non-gonadal sites, WAT being the main source...
of estrogen biosynthesis. However, it is not known how well blood hormone concentrations reflect the actual local synthesis of estrogens in the breast tissue. Recent studies indicate that postmenopausal women with normal BMI may exhibit signs of WAT inflammation. Furthermore, it should be noted that postmenopausal status is associated with increased expression of aromatase in BAT, independent of BMI. Therefore, it is not only the obese women who are at high risk of breast cancer, but also it is well possible that (some) postmenopausal women with normal BMI and blood estrogen levels may present with inflammation and increased aromatase expression in breast tissue, causing an increase in their cancer risk.

Taken together, cessation of ovarian hormone production causes a wide range of changes that lead to increase aromatase expression in BAT. Postmenopausal hormone milieu predisposes women to weight gain, as well as to BAT inflammation, which, eventually, may increase local estrogen biosynthesis and promote breast carcinogenesis. BAT, breast adipose fibroblast

FIGURE 6 Regulation of aromatase expression in the postmenopausal breast adipose stroma. Cessation of ovarian hormone production causes a wide range of changes that lead to increase aromatase expression in BAT. Postmenopausal hormone milieu predisposes women to weight gain, as well as to BAT inflammation, which, eventually, may increase local estrogen biosynthesis and promote breast carcinogenesis. BAT, breast adipose fibroblast

| Body weight | Normal BMI | Normal BMI | Obesity |
|-------------|------------|------------|---------|
| Adiposity   | Low-BMI    | Menopause-related increase | Increased |
| Status of the mammary parenchyma | Functional | Involved | Involved |
| Expression of aromatase (CYP19A1) | BAF, low-normal tissue levels | BAF, menopause-related increase | BAF, increased (up to 4-fold change) |
| CYP19A1-PLA1 transcript | + | + | ++++ |
| CYP19A1-PLA1 transcript | Very low | Very low | ++++ |
| Inflammation | No | Menopause-related increase | Low-grade inflammation | Increased macrophage infiltration |
| Representation of CYP19A1-PLA1 (II-11) | Normal | Low | Low |
| Inducers of CYP19A1-PLA1 (TNF-α, IL-1β, IL-6, IL-12) | Low | + | ++++ |
| Inducers of CYP19A1-PLA1 (PO22) | Very low | Very low | ++++ |

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CONFLICT OF INTEREST
The authors declare there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

AUTHOR CONTRIBUTION
GM, EY, NS, and SM designed the experiments and analyzed the data, EP and PH collected breast tissue samples and isolated primary breast ASCs, and GM, EY, LP, NS, and DD performed experiments. All authors participated in the preparation of the manuscript.

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SUPPORTING INFORMATION

Additional Supporting Information may be found online in the Supporting Information section.

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