In vitro maturation impacts cumulus–oocyte complex metabolism and stress in cattle

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Abstract

The influence of *in vitro* maturation (IVM) in oocytes is still not totally understood. The aim of this study was to determine the influence of IVM on the metabolism and homeostasis of bovine cumulus-oocyte complexes. In the present study, we demonstrated that IVM leads to accumulation of neutral lipids associated with differential levels of the mono-, di- and triacylglycerols in both cumulus cells and oocytes. We observed that *in vitro*-matured oocytes exhibited decreased glutathione and reactive oxygen species levels and a lower ATP/ADP ratio when compared to *in vivo*-matured oocytes, with no significant differences in metabolism and stress-related mRNA or miRNA levels. Moreover, in addition to an increase in lipids in *in vitro*-matured cumulus cells, fatty acid synthesis and accumulation as well as glycolysis pathway genes were upregulated, whereas those affiliated with the β -oxidation pathway were decreased. Our gene expression data in cumulus cells suggest the disruption of endoplasmic reticulum stress, apoptosis and cellular stress response pathways during IVM. Furthermore, a total of 19 miRNAs were significantly altered by the maturation process in cumulus cells. These results indicate some new negative influences of the *in vitro* system in cumulus-oocyte complexes, demonstrating the occurrence of functional disruption in lipid metabolism and stress pathways and showing evidences suggesting the occurrence of altered mitochondrial activity and energy metabolism during IVM, with a massive dysregulation of the corresponding transcripts in the surrounding cumulus cells.

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Introduction

Assisted reproduction techniques are widely used as a means to rescue fertility in several species and as a tool of genetic improvement in cattle. The *in vitro* system attempts to mimic the *in vivo* environment; however, there is a myriad of factors that can contribute to lower embryo production and quality. Although the culture environment is known to lead to stress and lipid accumulation after *in vitro* embryo production, impairing embryo cryotolerance and pregnancy rates in bovines (Goto *et al.* 1993, Wright & Ellington 1995, Abe *et al.* 2002, Rizos *et al.* 2002, Seidel 2006), it is unknown whether these effects are the consequence of exposure of the oocyte to the *in vitro* environment or if other underlying mechanisms may be involved. Therefore, it is important to perform comprehensive studies regarding these contributing factors during

oocyte *in vitro* maturation (IVM) for comparison with the *in vivo*-produced counterparts.

In this context, oocyte maturation is dependent on an orchestrated sequence of events involving oocyte and follicular constituents, such as cumulus cells, to produce oocytes that are competent to support development. Thus, studies considering cumulus-oocyte complexes (COCs) as interdependent compartments and focusing on their behavior during maturation are necessary to understand the influence of IVM on cell metabolism and homeostasis.

Oxidative phosphorylation constitutes the most important energy pathway during oocyte maturation. Given that oocytes and embryos are not able to perform glycolysis prior to the compaction and blastulation stages, the cumulus cells are responsible for providing pyruvate and lactate to the oocyte (Khurana & Niemann 2000, Thompson *et al.* 2015). However, β -oxidation, another metabolism pathway, has also been recognized as being

highly important during oocyte maturation. Studies have shown an increase of oocyte quality and embryo development rate when β -oxidation was increased during IVM, suggesting the occurrence of an energy imbalance during IVM and pointing to deficient β -oxidation as being likely responsible (Downs *et al.* 2009, Dunning *et al.* 2010, 2014, Paczkowski *et al.* 2013). Moreover, it was demonstrated that IVM leads to an increase in lipid accumulation that does not occur during *in vivo* maturation (Del Collado *et al.* 2016), suggesting an alteration in lipid metabolism during *in vitro* culture.

Furthermore, the properties of energy metabolism and its association with cellular stress responses during oocyte maturation also remain poorly understood. Although most studies consider metabolism and homeostasis modifications as independent effects during the *in vitro* process, some have demonstrated a correlation between increased lipids and cellular stress (Borradaile *et al.* 2006, Diakogiannaki *et al.* 2008).

Recently, the roles of certain miRNAs in maturation have been described, the majority of which are expressed in cumulus cells, although several effects were noted on oocyte maturation and embryo development as well (Hossain *et al.* 2012, Miles *et al.* 2012, Shi *et al.* 2015). In addition, the miRNA-mediated regulation of metabolism and stress-related genes has been demonstrated in several species (Duru *et al.* 2015, Yang *et al.* 2015, Magenta *et al.* 2016); however, it is still unknown how miRNAs modulate these genes during oocyte maturation in bovines.

Therefore, the aim of this study was to determine the mechanisms influencing cumulus cell and oocyte metabolism and homeostasis during IVM by analyzing energy substrates and metabolites as well as stress-related compounds, in addition to transcripts and post-transcriptional regulators of energy metabolism, ER stress, apoptosis and cellular responses to stress. We hypothesized that the *in vitro* environment causes metabolic disruption, increasing lipid accumulation, altering the mono-, di- and triacylglycerols profile, leading to cellular stress of the COC. In order to address these questions, we performed several measurements in oocytes, indicative of potential metabolic and stress-related modifications acquired during IVM. As we did not observe alterations in transcriptional or post-transcriptional regulators in oocytes that could explain the effects of the *in vitro* system, we investigated the mechanisms leading to the noted alterations in cumulus cells.

Materials and methods

Reagents were purchased from Sigma Chemical Co. unless otherwise indicated. The study was approved by the Bioethical Committee of the FZEA – University of Sao Paulo, Pirassununga, SP, Brazil, under protocol number 14.1.675.74.7. Experiments were conducted in accordance with International Guiding Principles for Biomedical Research

Involving Animals, meeting the Society for the Study of Reproduction standards.

Immature and *in vitro*-matured oocyte collection

We collected grade I and II COCs by postmortem follicular aspiration of 3–8 mm follicles from Nellore (*Bos indicus*) ovaries obtained from slaughterhouse carcasses to obtain immature and *in vitro*-matured oocytes. The denuded oocytes and corresponding cumulus cells were recovered from immature COCs and stored for further analysis. A fraction of the immature COCs was assigned for IVM at 38.5°C and 5% CO₂ in air and high humidity for 24 h. The IVM media was composed of TCM-199 with Earle's Salts, L-glutamine and 2.2 g/L sodium bicarbonate (GIBCO BRL), 8.75×10^{-4} IU/mL (0.5 µg/mL) follicle-stimulating hormone (FSH; Ourofino Saude Animal, Cravinhos, Brazil), 50 µg/mL human chorionic gonadotropin (hCG; Vetecor; Ourofino Saude Animal), 50 µg/mL gentamicin, 0.2 mM sodium pyruvate and 10% fetal bovine serum (FBS, GIBCO). After maturation, cumulus cells from expanded COCs were separated from the oocyte with needles. The oocytes were totally denuded by continuous pipetting with 135-µm-diameter stripper tips. All oocytes were evaluated for nuclear maturation and only MII oocytes presenting the first polar body (1 PB) and the corresponding cumulus cells were used for further analysis. Samples assigned for RNA were immediately placed in liquid nitrogen and stored at –80°C until further use. The samples from 12 different IVM routines were obtained simultaneously with *in vivo* oocyte collections.

In vivo-matured oocyte collection

To obtain *in vivo*-matured oocytes, the ovarian follicular wave was synchronized in 33 cyclic Nellore cows and animals were submitted to follicular superstimulation during 12 different ovum pick-up (OPU) sessions. On the first day (Day 0), females received 0.5 mg cloprostenol (Sincrocio, Ourofino Saude Animal) and follicular wave synchronization was induced by follicular ablation associated with the administration of 2 mg benzoate estradiol (Sincrodiol, Ourofino Saude Animal) and intravaginal progesterone-releasing device (IPRD) implants (Sincrogest, Ourofino Saude Animal). Follicular superstimulation was initiated on Day 4. A total of 133 mg of FSH (Follitropin, Bioniche Animal Health Canada Inc., Belleville, ON, Canada) per cow was divided into 8 decreasing doses administered every 12 h. Along with the sixth dose of FSH, 0.5 mg of cloprostenol was administered and 12 h later, the IPRD was removed. On Day 8, females received 0.02 mg buserelin acetate (GnRH analog, Sincroforte, Ourofino Saude Animal) and 25–26 h later expanded COCs were recovered using the OPU method. *In vivo*-matured oocytes were evaluated for nuclear maturation and selected based on 1 PB extrusion and only matured oocytes (identified as MII oocytes) and corresponding cumulus cells were used for further analysis.

High-performance liquid chromatography mass spectrometry analysis (HPLC–MS)

HPLC–MS was used to identify lipids in immature, *in vivo*- and *in vitro*-matured oocytes and cumulus cells (Supplementary

Fig. 1, see section on [Supplementary Data](#) given at the end of this article). For each group, five pools of cumulus cells retrieved from seven COCs each or five pools of ten denuded oocytes were used.

Chemicals and reagents

Methanol (MeOH, HPLC grade), chloroform (CHCl₃, HPLC), acetonitrile (ACN, HPLC), water (H₂O, ultra-gradient grade) and isopropanol (IPA, HPLC) were purchased from JT Baker (Avantor, Center Valley, PA, USA). MS reference standard d₅-Triacylglycerol Internal Standard mixture and the reference standard d₅-Diacylglycerol Internal Standard mixture were from Avanti Polar Lipids, Alabaster, AL, USA. Strata silica columns (100 mg for 1 mL) were from Phenomenex (Torrance, CA, USA).

Preparation of standard internal solutions

Reference substances were combined at 200 ng/mL each in CHCl₃:MeOH (1:1, v/v) and diluted to 40 ng/mL in ACN as a working solution.

Extraction of sample lipids

For each pool (10 µL in PBS buffer), 150 µL of water, 450 µL IPA and 50 µL internal standard mix (40 ng/mL) was added and vortexed. Then, the solution was sonicated (20 min), centrifuged (15,000 g for 5 min at room temperature) and the supernatant was collected and dried under a constant stream of nitrogen (45°C).

Light petroleum–diethyl ether (2 mL, 90:10, v/v) was added to the dried extract, which was cleaned-up with an SPE column (100 mg), previously conditioned with 2 mL of water and 5 mL of light petroleum–diethyl ether (90:10, v/v). The clean-up with the SPE column is important to remove phospholipids that could interfere with mono-, di- and triacylglycerol (MG, DG, and TG, respectively) measurements. Lipids were eluted with 5 mL diethyl ether containing MG, DG and TG (MárquezRuiz *et al.* 1996) and evaporated under a constant stream of nitrogen (45°C). Lipid extracts were diluted with 50 µL ACN:IPA:H₂O (79:5:1 – 5 mM ammonium acetate), vortexed (3 min), filtered with polyvinylidene fluoride membrane filters (0.22-µm pores) for 5 min (10,000 g) and injected to the HPLC. The identified lipids were quantified against their corresponding internal standard (for MG, TG was used as the internal standard).

Analysis

Lipids were quantified using a 5500 Q-TRAP mass spectrometer (Sciex, Framingham, MA, USA) with an ESI interface in positive mode and a 1260 series HPLC Agilent device (Agilent). Data acquisition and analyses were performed using Analyst software (version 1.6.1; Sciex). Labeled and unlabeled lipids were separated on a Phenomenex C8 column (3.0 × 150 mm, 2.7 µm, Phenomenex) at a flow rate of 300 µL/min. Mobile phase B (ACN:IPA 80:20, v/v) was linearly increased from 1 to 99% over 5 min, kept constant for 20 min, returned to the initial condition over 1 min and kept constant for an additional 5 min. Mobile phase A consisted of ACN:5 mM ammonium

acetate 1:1. Injection volume was 5 µL. The optimized ESI interface parameters were the following: capillary voltage, 5 kV; curtain gas 10 psi; nebulizer gas 45 psi; desolvation gas 20 psi; CAD gas 12 psi; source temperature, 450°C. [Supplementary Table 1](#) for optimized multiple reaction monitoring (MRM) parameters such as declustering potential, collision energies and collision cell exit potential for each lipid (synthetic lipid-labeled solutions).

Quantification of reduced glutathione (GSH), reactive oxygen species (ROS), ATP and ADP levels in immature and in vivo- and in vitro-matured oocytes

For ROS and GSH quantification, a total of 37 (immature), 29 (*in vivo*-matured) or 24 (*in vitro*-matured) oocytes were stained with 5 µM CellRoX Green (Molecular Probes) and 10 µM Celltracker Blue CMF2HC (Molecular Probes) and fixed according to the manufacturer's instructions. Then, the oocytes were analyzed on an LSM 710 confocal microscope (Carl Zeiss) using a 63× objective in oil and an Argon laser (488 nm) and diode (405 nm), with the excitation/emission adjusted for each probe parameter. All images were captured under the same parameters, using sequential acquisition. We captured three images of each oocyte; these were assessed using ImageJ software (NIH; <http://rsb.info.nih.gov/ij/>) to measure the fluorescence intensity (f.i.) of GSH and ROS.

ATP and ADP content and the ATP/ADP ratio of 30 individual oocytes per group in five replicates were measured using the bioluminescence ApoSENSOR ATP Assay Kit (BioVision) according to the manufacturer's instructions. Briefly, we measured the ATP absorbance by a luminescence analysis. Then, the samples underwent an enzymatic reaction to convert the ADP to ATP, and the ATP absorbance was measured again. The levels of ADP were calculated from the difference in absorbance values obtained in the first and second measurements. A five-point ATP standard curve (0.15, 0.075, 0.0375, 0.01875 and 0.009375 µM) was used to estimate sample concentration.

RNA extraction and messenger RNA and microRNA reverse transcription

For total RNA extraction, we used five pools of 20 immature and *in vivo*- or *in vitro*-matured oocytes and five or eight different cumulus cell pools retrieved from 20 immature or 10 *in vivo*- and *in vitro*-matured COCs. Total RNA extraction was performed using TRIzol reagent and miRNAse kit (Qiagen). Reverse transcription of mRNA was performed using the High-Capacity cDNA Reverse Transcription kit (Applied Biosystems); for miRNA reverse transcription, we utilized the miScript polymerase chain reaction (PCR) system (Qiagen #218193) according to manufacturer instruction.

Real-time PCR analysis of genes related to lipid metabolism and to cellular stress in immature, in vivo- and in vitro-matured oocytes and cumulus cells

For mRNA expression, we performed microfluidic transcript level analysis of 25 cellular metabolism and stress-related

genes (Supplementary Table 2) utilizing the Biomark HD System (Fluidigm, South San Francisco, CA, USA). For this analysis, we used *PPIA*, *GAPDH* and *ACTB* as endogenous control genes. Microfluidic gene expression analysis was performed using *Bos taurus*-specific TaqMan assays (20×, Applied Biosystems). After the pre-amplification process using TaqMan PreAmp Master Mix (Applied Biosystems), samples were diluted 5-fold and analyzed by reverse transcription-quantitative PCR (RT-qPCR) using the TaqMan Universal PCR Master Mix (2×, Applied Biosystems) and TaqMan assays (20×, Applied Biosystems) in the 96.96 Dynamic Array Integrated Fluidic Circuits on the Biomark HD System (Fluidigm) using the protocol TaqMan GE 96×96 Standard, according to manufacturer's instructions.

To complement this work, we also performed real-time RT-qPCR of several genes related to lipid synthesis and accumulation and transport (*PLIN2*, *PLIN3* and *SREBP1*), β -oxidation (*CPT1A*, *CPT1B* and *CPT2*) and endoplasmic reticulum (ER) stress (*CHOP10*, *IRE*, *PERK*, and *ATF6*), where *PPIA*, *SDHA* and *YWHAZ* were used for the endogenous control genes. Primers were designed based on bovine sequences available at GenBank (Supplementary Table 3); these were tested for efficiency and the obtained amplicons were sequenced to ensure their specificity. Real-time RT-qPCR reactions were performed using 10 ng cDNA for each gene and 75 nM of each primer with the Power SYBR Green PCR Master Mix kit (Applied Biosystems) in the QuantStudio 6 Flex PCR System (Applied Biosystems).

Real-time PCR expression analysis of miRNAs predicted to regulate genes involved in cellular metabolism and stress in immature, *in vivo*- and *in vitro*-matured oocytes and cumulus cells

To evaluate whether miRNAs predicted to regulate the studied genes were altered following IVM, we selected miRNAs identified using bioinformatics. The relative levels of 88 bovine mature miRNAs (Table 1 and Supplementary Table 4) were examined in immature, *in vivo*- and *in vitro*-matured oocytes and cumulus cells. Specifically, the miRNAs evaluated in this study were chosen based on validated or predicted interaction with the target genes. We used three criteria for miRNA inclusion. First, miRNAs already validated for bovine genes using miRTarBase (<http://mirtarbase.mbc.nctu.edu.tw/>) were directly included in this study. Next, those without validation for bovine genes were compared with their homologs validated for humans by analyzing the similarity between the human and bovine miRNAs using miRBase (<http://mirbase.com>, Supplementary Table 5) and their binding site on the target gene as predicted by TargetScan software (www.targetscan.org). Finally, if the target gene had no validated miRNA, we chose bovine miRNAs predicted to target the genes using TargetScan.

The RT-qPCR reaction was performed using Quantitect SYBR Green (Qiagen), 10 μ M Universal reverse primer (Qiagen), 10 μ M miRNA-specific forward primer (Supplementary Table 6), and 60 pg cDNA and analyzed using the QuantStudio 6 Flex PCR System. The Ct values were normalized with the geometric mean

Table 1 Bovine miRNA amplified in this study and their respective target genes.

miRNAs	Target genes
bta-let-7a-5p	<i>CASP3</i>
bta-miR-15a	<i>ACSL6</i> , <i>BCL2</i>
bta-miR-15b	<i>FASN</i> , <i>ACSL6</i> , <i>BCL2</i>
bta-miR-16a	<i>ACSL6</i> , <i>BCL2</i>
bta-miR-16b	<i>FASN</i> , <i>ACSL6</i> , <i>BCL2</i>
bta-miR-17-5p	<i>PFKP</i> , <i>CASP9</i>
bta-miR-20a	<i>PFKP</i> , <i>CASP9</i>
bta-miR-20b	<i>PFKP</i> , <i>CASP9</i>
bta-miR-23a	<i>ACSL3</i>
bta-miR-23b-3p	<i>ACSL3</i>
bta-miR-24-3p	<i>ACSL6</i>
bta-miR-25	<i>ATF6</i>
bta-miR-26a	<i>ACSL3</i>
bta-miR-26b	<i>ACSL3</i>
bta-miR-27a-3p	<i>NRF2</i>
bta-miR-29b	<i>BAX</i>
bta-miR-29c	<i>BAX</i>
bta-miR-29d-3p	<i>BAX</i>
bta-miR-30a-5p	<i>GRP78</i>
bta-miR-30b-5p	<i>CAT</i>
bta-miR-30c	<i>CASP3</i>
bta-miR-30d	<i>CASP3</i>
bta-miR-32	<i>ATF6</i>
bta-miR-33a	<i>SREBP1</i> , <i>CPT1A</i>
bta-miR-33b	<i>CPT1A</i>
bta-miR-34a	<i>BCL2</i>
bta-miR-34b	<i>BCL2</i>
bta-miR-34c	<i>BCL2</i>
bta-miR-92a	<i>ATF6</i>
bta-miR-92b	<i>ATF6</i>
bta-miR-93	<i>PFKP</i> , <i>CASP9</i>
bta-miR-96	<i>CHOP10</i>
bta-miR-106a	<i>PFKP</i> , <i>CASP9</i>
bta-miR-106b	<i>PFKP</i> , <i>CASP9</i>
bta-miR-124a	<i>CPT1A</i>
bta-miR-124b	<i>CPT1A</i>
bta-miR-125a	<i>ATF6</i>
bta-miR-125b	<i>BCL2</i>
bta-miR-132	<i>ACSL3</i>
bta-miR-136	<i>SOD2</i>
bta-miR-137	<i>ACSL6</i>
bta-miR-142-5p	<i>ACSL6</i> , <i>CASP9</i> , <i>NRF2</i>
bta-miR-143	<i>PGK1</i> , <i>BCL2</i>
bta-miR-144	<i>NRF2</i>
bta-miR-146a	<i>PGK1</i>
bta-miR-146b	<i>PGK1</i>
bta-miR-149-5p	<i>ATF6</i>
bta-miR-153	<i>NRF2</i>
bta-miR-181a	<i>SCD</i> , <i>ACSL6</i> , <i>CPT1A</i> , <i>BCL2</i>
bta-miR-181b	<i>ACSL6</i> , <i>CPT1A</i> , <i>BCL2</i> , <i>GRP78</i>
bta-miR-181c	<i>ACSL6</i> , <i>CPT1A</i> , <i>BCL2</i>
bta-miR-181d	<i>ACSL6</i> , <i>CPT1A</i> , <i>BCL2</i>
bta-miR-182	<i>BCL2</i>
bta-miR-186	<i>ACSL6</i> , <i>CASP6</i>
bta-miR-195	<i>ACSL6</i> , <i>BCL2</i>
bta-miR-199a-3p	<i>ATF6</i>
bta-miR-199a-5p	<i>IRE1</i> , <i>ATF6</i> , <i>GRP78</i>
bta-miR-200b	<i>ACSL3</i>
bta-miR-200c	<i>ACSL3</i>
bta-miR-202	<i>ACSL3</i>
bta-miR-204	<i>ELOVL6</i> , <i>BCL2</i>
bta-miR-206	<i>G6PD</i>

bta-miR-211	<i>ELOVL6, BCL2</i>
bta-miR-218	<i>ACSL6</i>
bta-miR-223	<i>ACSL3</i>
bta-miR-302b	<i>PFKP</i>
bta-miR-302c	<i>PFKP</i>
bta-miR-302d	<i>PFKP</i>
bta-miR-330	<i>ACSL6, SOD2</i>
bta-miR-346	<i>CASP9</i>
bta-miR-365-3p	<i>PGK1, BCL2</i>
bta-miR-374a	<i>ACSL6</i>
bta-miR-374b	<i>ACSL6</i>
bta-miR-378	<i>ATF6</i>
bta-miR-381	<i>ACSL3</i>
bta-miR-421	<i>ACSL3</i>
bta-miR-424-5p	<i>ACSL6</i>
bta-miR-429	<i>ACSL3, CPT2</i>
bta-miR-448	<i>ACSL6</i>
bta-miR-449a	<i>BCL2</i>
bta-miR-495	<i>ACSL6, PGK1</i>
bta-miR-497	<i>ACSL6, BCL2</i>
bta-miR-499	<i>CPT1A</i>
bta-miR-503-5p	<i>ACSL6, ATF6</i>
bta-miR-582	<i>ACSL6</i>
bta-miR-670	<i>ATF6</i>
bta-miR-873	<i>PGK1, GPX1</i>
bta-miR-875	<i>ATF6</i>

of *RNU43* snoRNA, Hm/Ms/Rt U1 snRNA and bta-miR-99b. The expression value was calculated using the method of $2^{-\Delta Ct}$.

Statistical analysis

The lipid accumulation and the ATP and ADP data were compared using the non-parametric Kruskal–Wallis test. Data of GSH and ROS quantification along with MG, DG and TG quantitation as well as the mRNA and miRNA expression data were subjected to analysis of variance (ANOVA) and the averages were compared using the Tukey test. The equality of the variances was tested using the Levene test and normality was assessed with the Shapiro–Wilk test. All analyses were performed using SAS software (SAS Institute Inc., Cary, NC, USA) and a level of significance of 5% was considered for all tests.

Results

In vitro culture of COCs increased DGs and TGs in oocytes

No difference was observed in the total amount of MGs in oocytes after IVM when compared to the *in vivo* system (Fig. 1A). However, when analyzing

the individual profiles of MGs, we noticed a general increase of long-chain fatty acids in *in vitro* vs *in vivo* oocytes (Supplementary Fig. 2A). In particular, seven MGs increased during both *in vivo* and *in vitro* maturation (12:0, 14:0, 16:0, 18:2, 18:3, 24:0 and 26:1; Supplementary Fig. 2A). Of these, only MG 26:1 was increased in *in vivo*-matured oocytes compared to *in vitro*, whereas MGs 12:0, 16:0 18:1, 18:3, 20:4 and 24:0 showed higher levels in IVM compared to *in vivo*-matured oocytes (Supplementary Fig. 2A). In addition, five MGs were increased solely after IVM (14:1, 16:1, 18:0, 18:1 and 20:0), whereas MG18:1 was decreased after *in vivo* maturation when compared to immature oocytes.

In comparison, the total levels of DGs and TGs increased only in the IVM-derived oocytes (Fig. 1B and C). We identified eight DGs that increased during both maturation systems ((16:0)₂, (17:0)₂, (19:0)₂, (20:2)₂, (20:5)₂, (24:0)₂, (24:1)₂ and (24:4)₂) and five that increased only in IVM ((18:0)₂, (18:2)₂, (18:3)₂, (20:4)₂ and (22:4)₂) (Supplementary Fig. 2B). Among those that increased during both maturation systems, four DGs showed a preferential increase in the *in vitro* compared to the *in vivo* group ((16:0)₂, (24:0)₂, (24:1)₂, and (24:4)₂). Additionally, with the exception of (17:0)₂ 17:1, which exhibited no difference between the *in vivo* and *in vitro* groups, all the other eight TGs species quantified in this study ((14:0)₂ 16:1, (15:0)₂ 18:1, (16:0)₂ 18:0, (16:2)₂ 18:1, (20:0)₂ 20:1, (20:2)₂ 18:2, (20:4)₂ 18:2 and (20:5)₂ 22:6) were higher in *in vitro*-matured oocytes compared to their *in vivo*-matured counterparts (Supplementary Fig. 2C).

IVM decreases GSH levels and ATP/ADP ratio in oocytes compared to *in vivo* maturation

GSH levels increased in both maturation processes compared to immature oocytes (15.43 ± 7.45 f.i.), although we observed relatively higher levels in *in vivo*- (30.49 ± 13.80 f.i.) compared to *in vitro*-matured oocytes (22.84 ± 8.90 f.i.) (Fig. 2A). Furthermore, ROS levels between immature (11.39 ± 4.30 f.i.) and matured groups did not differ (Fig. 2B). However, ROS levels were higher in the *in vivo* (13.53 ± 4.54 f.i.) compared with the *in vitro* (10.72 ± 3.26 f.i.) group (Fig. 2B).

We also observed increased ATP levels in the *in vitro* (1.784 ± 0.333 pmol/oocyte) compared to the

Quantification of neutral lipids in oocytes by HPLC-MS/MS

Figure 1 Quantitation of neutral lipids in oocytes by HPLC–MS/MS. Absolute quantities of monoacylglycerols (A), diacylglycerols (B), and triacylglycerols (C) in oocytes from immature, *in vivo*-matured, and *in vitro*-matured COCs. Different letters above the bars within a graph indicate significant differences ($P < 0.05$). Bars depict the means and error bars depict S.E.M.

Figure 2 GSH (A) and ROS (B) quantification before (immature) and after *in vivo* and *in vitro* maturation in oocytes. Different letters above the bars for the same graph indicate significant differences ($P < 0.05$). Bars depict the means and error bars depict S.E.M.

in vivo (1.551 ± 0.420 pmol/oocyte) and immature (1.437 ± 0.672 pmol/oocyte) groups. Additionally, we observed decreased ADP levels in *in vivo* oocytes (0.127 ± 0.126 pmol/oocyte) compared to *in vitro* (0.235 ± 0.165 pmol/oocyte) and immature (0.176 ± 0.110 pmol/oocyte) oocytes ($P = 0.06$; Fig. 3A and B respectively for ATP and ADP). Furthermore, to evaluate the energy demand, we calculated the ATP/ADP ratio. The *in vivo* group (36.88 ± 37.34) showed an increased energy demand when compared to *in vitro* (12.87 ± 12.53) and immature (12.74 ± 12.73) groups (Fig. 3C).

Figure 3 Scattered dot plots for ATP (A), ADP (B), and ATP/ADP (C) ratio quantities in immature and *in vivo*- and *in vitro*-matured oocytes. Each dot represents an oocyte, lines indicate the means, and brackets indicate statistical difference ($P < 0.05$; unless differently stated) between groups.

Cellular metabolism and stress-related genes are not altered in *in vitro*-matured oocytes

With the exception of the *PGK1* gene, we did not detect the statistical differences in the levels of the selected genes among the oocyte groups (Supplementary Fig. 3 and Supplementary Table 7).

The majority of miRNAs exhibited similar levels in oocytes after maturation *in vivo* or *in vitro*

Only six out of 88 miRNAs were altered in oocytes after maturation (Fig. 4). Analysis of oocytes showed decreased levels of miR-17-5p after IVM when compared to immature oocytes, whereas six miRNAs (let-7a-5p, miR-106a, miR-106b, miR-17-5p, miR-23b-3p and miR-365-3p) were significantly decreased in *in vivo*-matured oocytes compared to immature oocytes. Notably, analysis of matured oocytes demonstrated two miRNAs (bta-miR-106b and bta-miR-17-5p) that were significantly decreased in *in vivo*-matured oocytes compared to *in vitro* oocytes. The results of the relative expression of all tested miRNAs in all groups are shown in Supplementary Table 8.

IVM of COCs increases TGs in cumulus cells

The total amount of MGs increased after maturation in the cumulus cells of both the *in vivo* and *in vitro* groups (Fig. 5A). Compared to the *in vivo* group, however, the *in vitro* cumulus cells exhibited a lower total amount of MGs, a pattern that was also followed by three specific MGs (12:0, 20:4, and 22:2). In contrast, six MGs (14:1, 16:1, 18:1, 18:3, 24:0 and 26:1) were higher in *in vitro* than *in vivo* cumulus cells. It is noteworthy that the total amount as well as eight specific MGs (12:0, 16:0, 16:1, 18:0, 18:1, 18:3, 24:0 and 26:1) increased both in *in vivo*- and *in vitro*-derived cumulus cells compared to their immature counterparts, whereas 14:1 increased only in IVM and 14:0, 18:2, 20:0, 20:4 and 22:2 increased solely during *in vivo* maturation (Supplementary Fig. 2D).

The total amount of TGs increased only after IVM in cumulus cells (Fig. 5C). Similarly, six ((15:0)₂ 18:1, (16:0)₂ 18:0, (16:2)₂ 18:1, (17:0)₂ 17:1, (20:0)₂ 20:1, (20:2)₂ 18:2 and (20:4)₂ 18:2) out of eight species of TGs were higher in the *in vitro* compared to the *in vivo* group (Supplementary Fig. 2F). All nine TG species accumulated during IVM in comparison with those in

Figure 4 Hierarchical clustering and heatmap depicting expression levels of miRNAs differentially expressed among immature, *in vivo*-matured, and *in vitro*-matured oocytes.

immature cumulus cells; however, this pattern was not observed for four TG species ((16:0)₂ 18:0, (16:2)₂ 18:1, (20:2)₂ 18:2 and (20:4)₂ 18:2) between immature and *in vivo* groups (Supplementary Fig. 2F).

Total DG levels were also higher in cumulus cells obtained after IVM compared to immature or *in vivo* controls. These data, as well as the levels of DG species are provided in Fig. 5C and Supplementary Fig. 2E.

Cellular metabolism and stress-related genes are altered in *in vitro*-matured cumulus cells

The relative levels of cellular metabolism-related genes in cumulus cells differed for 27 out of 34 genes during maturation (Fig. 6 and Supplementary Table 7); additionally, a total of sixteen genes showed differences between *in vivo* and *in vitro* groups (Fig. 6). The metabolism-related genes *ACACA*, *PLIN2*, *SCD*, *ACSL3*, *CPT1A*, *CPT2*, *G6PD* and *PKG1* presented increased levels during maturation regardless of the system when compared to the immature group (Fig. 6). Of these, *ACACA*, *PLIN2*, *SCD*, *G6PD* and *PKG1* transcripts exhibited relatively higher levels in IVM. However, *CPT1A* demonstrated higher levels in *in vivo*, when compared to cumulus cells from *in vitro*-matured COCs. Conversely, *SREBP1*, *FASN* and *ELOVL6* only exhibited decreased expression levels during *in vivo* maturation (Fig. 6), whereas *SREBP1*, *ACSL6*, *CPT1B* and *PFKP* transcript levels were increased only during IVM.

ER stress and apoptosis-related genes such as *ATF6*, *BAX*, *BCL2* and *CASP9* were increased during maturation, whereas *CHOP10* and *GRP78* were decreased in both maturation groups (Fig. 6). Of these, *BAX* and *BCL2*

expression was relatively higher in cumulus cells from the *in vitro* compared with the *in vivo* group. Additionally, *CASP3* levels were increased only during IVM, whereas *IRE1*, *PERK* and *ATF4* transcripts maintained stable levels during maturation. The ER stress response and antioxidant-related genes *NRF2*, *KEAP1*, *SOD1*, *GPX4*, *PRDX1* and *CAT* patterns also changed during maturation. *NRF2* as well as the gene for its inhibitor, *KEAP1*, increased during both maturation systems, although the *in vivo* group demonstrated relatively higher *NRF2* transcript levels when compared to the IVM group. Conversely, *GPX4* transcript levels decreased regardless of the maturation system. Moreover, *SOD* and *CAT1* were increased in *in vitro*-matured cumulus cell compared to immature, while *PRDX1* was higher in *in vivo*-matured cumulus cell compared to immature and *in vitro*-matured cumulus cells.

MicroRNA levels are altered in cumulus cells after maturation *in vitro* and *in vivo*

A total of 19 miRNAs varied owing to the maturation process (Fig. 7). Of these, eight miRNAs (miR-181b, miR-181d, miR-182, miR-204, miR-218, miR-27a-3p, miR-29d-5 and bta-mir-96) significantly increased and two miRNAs (miR-23a and miR-23b-3p) significantly decreased in cumulus cells following IVM compared to those from immature follicles. In comparison, 11 miRNAs (miR-181d, miR-182, miR-186, miR-199a-3p, miR-200c, miR-223, miR-27a-3p, miR-29c, miR-29d-5p, miR-424-5p and miR-96) were significantly increased and three miRNAs (miR-17-5p, miR-23b-3p, and miR-330) were significantly decreased in cumulus cells after *in vivo* maturation compared to those from immature follicles. Notably, two miRNAs (miR-218 and miR-330) were significantly increased and six miRNAs (miR-182, miR-199a-3p, miR-200c, miR-223, miR-23a and miR-424-5p) were significantly decreased in cumulus cells after *in vitro* compared to *in vivo* maturation. The expression values of all tested miRNAs from all cell groups are listed in Supplementary Table 8.

Discussion

Despite all efforts to understand how the *in vitro* system influences oocyte and embryo development, limitations

Figure 5 Quantitation of neutral lipids in cumulus cells (CCs) by HPLC-MS/MS. Absolute quantities of monoacylglycerols (A), diacylglycerols (B), and triacylglycerols (C) in cumulus cells from immature, *in vivo*-matured, and *in vitro*-matured COCs. Different letters above the bars within a graph indicate significant differences ($P < 0.05$, except for letters 'a' and 'b' in A, which indicate $P = 0.06$). Bars depict the means and error bars depict s.e.m.

Figure 6 Expression of cellular metabolism and stress-related genes in cumulus cells from immature and *in vivo*- and *in vitro*-matured COCs. Different letters above the bars for the same gene indicate significant differences ($P < 0.05$). Bars depict the means and error bars depict S.E.M.

remain that render IVM far from equivalent to the *in vivo* conditions. Nevertheless, the means by which the oocyte maturation environment might influence these features are not clearly understood. Accordingly, in the current study, we addressed how the *in vitro* system influences metabolism and cellular homeostasis during oocyte maturation.

In particular, although some studies have described a decrease in lipid quantity in oocytes during IVM (Ferguson & Leese 1999, Kim *et al.* 2001), we demonstrated by MS analysis that, unlike *in vivo* maturation, IVM leads to increased DGs and TGs in oocytes and cumulus cells. This result is in accordance to other studies that demonstrated higher lipid accumulation in oocytes after IVM, even without serum supplementation (Aardema *et al.* 2011, Del Collado *et al.* 2016). In addition, we demonstrated that IVM could modify MG, DG and TG patterns in COCs. Notably, different fatty acids may be helpful or harmful

during maturation (McKeegan & Sturmeay 2011), indicating a possible influence of some fatty acid on oocyte quality during IVM. For example, palmitic acid (16:0) and linolenic acid (18:3) were showed to have detrimental effects in oocyte maturation and embryo development upon their appearance in high concentration during IVM (Leroy *et al.* 2005, Marei *et al.* 2009). As in our results, both fatty acids increased in oocytes in *in vitro*-matured compared with *in vivo*-matured groups, which indicates higher levels than what would be expected in physiological conditions. Detrimental effects owing to the levels of these fatty acids could occur during IVM. However, the exact roles and biological significance of each specific lipid species in COCs maturation or their ability to develop remain largely unknown.

Furthermore, certain lipid synthesis and accumulation genes were increased in *in vitro*-matured cumulus cells, demonstrating that these pathways were upregulated

Figure 7 Hierarchical clustering and heat map depicting the expression levels of miRNAs differentially expressed among immature, *in vivo*-matured, and *in vitro*-matured cumulus cells.

compared to cumulus cells from *in vivo*-matured COCs. Our data confirmed our hypothesis that, during IVM, lipid synthesis occurred in cumulus cells. Additionally, fatty acid desaturation and elongation genes were also influenced, and these results were confirmed by MS analysis, from which we observed that the majority of the long-chain desaturated fatty acids were identified as increased in cumulus cells after *in vitro* culture.

Furthermore, our results suggest possible effects in mitochondrial activity and energy metabolism. The expression of β -oxidation and glycolysis-related genes in cumulus cells and ATP/ADP ratio in oocytes are indicative of reduced β -oxidation and increased glycolysis during IVM within COC, whereas the results of the ATP/ADP ratio demonstrated an alteration in energy demand in *in vitro* oocytes when compared to their *in vivo* counterparts. We note, however, that a wide spread of values was shown in *in vivo*-matured oocytes. Conversely, the higher similarity among *in vitro* samples, not observed in the *in vivo* group, likely occurred owing to the more similar conditions imposed to COCs during grouped *in vitro* culture. In addition to our ATP/ADP results in oocytes, cumulus cell gene expression data were in accordance with this metabolic behavior, wherein the glucose metabolism genes *G6PD*, *PFKP* and *PGK1* were increased and the most highly expressed β -oxidation *CPT1* gene, *CPT1A*, was decreased in the *in vitro* groups. An increase of the glycolysis pathway during IVM therefore may be a consequence of β -oxidation deficiency, providing an alternative form of energy supply. However, differences in glucose metabolism between *in vivo*- and *in vitro*-matured oocytes and cumulus cells are yet to be demonstrated.

In particular, the differential behavior of *CPT1A* and *CPT1B* transcript levels between *in vivo* and *in vitro*

systems potentially indicates that pre-transcriptional factors may influence these genes during IVM. However, we did not observe differences in DNA methylation as assessed by bisulfite DNA-sequencing analysis of the *CPT1A* promoter region of *in vitro*- and *in vivo*-matured cumulus cells (detailed in [Supplementary data 1](#)), suggesting that other mechanisms are likely involved in the transcriptional regulation of this gene.

Therefore, alterations in energy metabolism pathways identified in this study could, in part, be involved in lipid accumulation during IVM. Other authors have demonstrated that *in vivo*-matured oocytes showed higher mitochondria activity ([Zeng et al. 2014](#)) and lower β -oxidation levels when compared to *in vivo*-matured ones ([Dunning et al. 2010, 2014](#)). Altogether, these results suggest that there is a decrease in β -oxidation during IVM, which could lead to an increase of lipids in COCs.

Considering that our results and the literature suggest an increase in mitochondria activity in the *in vivo*-matured oocytes, the increase of ROS levels observed in this group was expected. However, the amount of GSH, which constitutes the most abundant antioxidant in oocytes and embryos, drastically increased during *in vivo* maturation when compared with IVM. Oocytes must establish the GSH pool during maturation because, after this stage, its synthesis is halted until subsequent activation of the embryo genome occurs ([Eichenlaub-Ritter et al. 2011](#)). This observation suggests an increased susceptibility to oxidative damage during *in vitro* embryo development.

Cumulus cells subjected to *in vitro* processes showed increased *BAX*, *BCL2* and *CASP3* levels when compared to *in vivo* maturation, indicating an upregulation of apoptosis pathways. Additionally, although we detected higher levels of *SOD1* in *in vitro*-matured cumulus cells, the levels of *NRF2* and *PRDX1* were lower compared to the *in vivo* process. Specifically, the *NRF2* transcription factor has been demonstrated to be crucial for activating the most important antioxidant pathway and is associated with survival and the developmental competence of embryos cultured under conditions of oxidative stress ([Amin et al. 2014](#)). Therefore, our results indicate that *in vitro* culture leads to cellular stress without enhancing the antioxidant pathway. These findings are in accordance with those observed by Tesfaye and coworkers who showed an increase in cellular stress-related genes in cumulus cells retrieved from an IVM system ([Tefsaye et al. 2009](#)).

To the best of our knowledge, this is the first study evaluating metabolism and stress-related miRNAs during *in vivo* and *in vitro* maturation. As only six out of 88 miRNAs investigated varied during maturation and only two, bta-miR-17-5p and bta-miR-106b, differed between *in vivo*- and *in vitro*-matured oocytes, the post-transcriptional regulation of these pathway by miRNAs did not appear to constitute the driving mechanism leading to the observed differences in oocyte metabolism.

The expression of mRNAs and miRNAs in cumulus cells showed both direct and indirect regulation of lipogenesis, with several components of these pathways varying between *in vivo*- and *in vitro*-derived COCs. In particular, the fatty acid elongation genes *ACSL3* and *ACSL6* exhibited different transcript levels after maturation. The miRNA-200c is predicted to regulate *JUN*, which leads to increased SREBP1 levels (Guo *et al.* 2016). Decreased expression levels of miR-200c in *in vitro*-matured cumulus cells may therefore lead to an increase in SREBP1 and, consequently, *FASN* and *TG* levels. Furthermore, miR-182 and miR-96, which were decreased in *in vitro*-matured cumulus cells, are predicted to regulate *FBW7* and *INSIG2*, thereby negatively regulating SREBP1 synthesis (Jeon *et al.* 2013). Although these mechanisms have been described in humans and mice, they may also be conserved in bovines according to TargetScan. Therefore, changes in the levels of miR-200c, miR-182 and miR-96 in the same cells may control SREBP1 synthesis, which in our study was increased in *in vitro* cumulus cells.

Our gene expression results during *in vivo* and *in vitro* maturation paralleled those in the literature. Despite major gene expression differences observed during embryonic gene activation between 4- and 8-cell embryos (Jiang *et al.* 2014), several authors have observed a difference in gene expression between oocytes from *in vitro* and *in vivo* maturation processes (Kues *et al.* 2008, Katz-Jaffe *et al.* 2009). Moreover, in *in vitro*-matured oocytes, genes related to the tricarboxylic acid cycle and oxidative phosphorylation were downregulated when compared to their expression in *in vivo*-matured oocytes, indicating a metabolic disruption during IVM (Katz-Jaffe *et al.* 2009). Moreover, as mentioned previously, Tesfaye and coworkers observed an increase of cellular stress genes in *in vitro*-matured cumulus cells (Tesfaye *et al.* 2009). However, we could not discard the possibility that superovulation treatment in *in vivo* samples may have interfered in some results. Several authors have shown that FSH treatment during superovulation influences miRNA expression (Noferesti *et al.* 2013) and gene expression (Chu *et al.* 2012, Fortier *et al.* 2014). Nevertheless, until now, an interference in metabolism and cellular stress gene expression after FSH treatment has not been described. Altogether, our work and the literature data confirm that *in vitro* culture alters metabolism and cellular stress pathways in COCs.

Lipid accumulation, primarily of long-chain fatty acids such as palmitate, causes detrimental effects on mitochondria and ER leading to ER stress and apoptosis (Borradaile *et al.* 2006, Diakogiannaki *et al.* 2008). Moreover, it has been showed that bovine, mouse and human oocytes with excessive lipid accumulation exhibit increased levels of ER stress markers (Wu *et al.* 2010, Yang *et al.* 2012). As we showed that IVM can lead to increased lipid quantities in COCs and that these

fatty acids primarily consist of a long-chain fatty acid, we may expect higher levels of stress in these cells. The results observed in the present study support that an increase in lipid accumulation consequent to the IVM system and a decrease of β -oxidation, combined with *in vitro* oxidative stress, may lead to the induction of ER stress and apoptosis.

Based on the negative relationship between lipids quantities, cell stress and apoptosis, it is important to understand the mechanisms involved in lipid accumulation during IVM. We hypothesized three possible pathways for lipid accumulation in the *in vitro* system: (1) an increase in lipids synthesis; (2) failure of β -oxidation and (3) fatty acid incorporation into oocytes. Here, we demonstrated a possibility of an increase of lipid synthesis and decrease of β -oxidation during IVM. Moreover, it was demonstrated that, depending on the type of supplementation (FBS or BSA), lipid accumulation occurred at different levels (Del Collado *et al.* 2016). We showed that the increased lipid accumulation in COCs is associated with a massive deregulation of lipid metabolism pathways in *in vitro*-matured cumulus cells, with no corresponding alterations in the gene expression profiles of the oocytes. Therefore, based on these results, it is tempting to speculate the existence of lipids trafficking to the oocyte cytoplasm, which may originate from the cell culture medium or from cumulus cells. We recently described potential fatty acid trafficking from cumulus cells to an oocyte via fatty acid-binding protein 3 and trans-zonal projections during oocyte maturation. This mechanism of lipid accumulation in oocytes is possibly more active during IVM when compared to *in vivo* maturation (Del Collado *et al.* 2017).

Thus, we demonstrated that IVM provided metabolically disrupted and stressed COCs. Furthermore, the *in vitro* system caused an increase in lipid synthesis and accumulation, modifications in the mono-, di- and triacylglycerols profiles, and an increase in cellular stress. Our results also suggest that IVM could affect mitochondrial activity, β -oxidation and glycolysis pathways during IVM, although more in-depth studies are needed to confirm this hypothesis. The mechanisms leading to these final cellular events are summarized in Fig. 8. In conclusion, our study provides evidence of metabolism and stress-related modifications during IVM of bovine oocytes. The results presented here encompass differences between *in vivo* and *in vitro*-matured COCs that may be attributed to the hormonal treatment prior to OPU, by the presence of serum in IVM or by several other potential factors influencing *in vivo* or *in vitro* maturation. The specific contribution of each of these factors is yet to be ascertained. These results highlight the necessity for further studies regarding the elucidation of cumulus cell–oocyte communication in order to better understand the underlying mechanisms involved in the poor quality of *in vitro*-produced oocytes and embryos.

Figure 8 Summary of the modifications caused by the *in vitro* system in metabolism and cellular stress in cumulus cells and oocytes. Herein, we summarize the most important results that support the hypothesis that *in vitro* maturation leads to increase of lipid synthesis and accumulation, causing cellular stress within cumulus-oocyte complexes: in cumulus cells, besides the increase of neutral lipids, the expression levels of mRNAs (*SREBP1*, *FASN* e *PLIN2*) and miRNAs (miR-96 and miR-182) suggested an increase of lipid synthesis and accumulation in *in vitro* matured cumulus cells. Moreover, we showed the increase of stress-related genes (*CASP3*, *BCL2* and *BAX*), the decrease of antioxidant pathway related gene (*NRF2*) and the increase of glycolysis-related genes (*G6PD*, *PFKP* and *PKG1*) in *in vitro* matured cumulus cells when compared to *in vivo* matured cumulus cells. In oocytes, we demonstrated the increase of lipids and ROS levels and the decrease of GSH levels and ATP/ADP ratio in *in vitro* matured oocytes when compared to *in vivo*-matured oocytes. LD, lipid droplet; TAC, tricarboxylic acid cycle; ETC, electron transfer chain; ARE, antioxidant response element; ER, endoplasmic reticulum; ROS, reactive oxygen species.

Supplementary data

This is linked to the online version of the paper at <http://dx.doi.org/10.1530/REP-17-0134>.

Declaration of interest

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

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