DIFFERENTIALLY EXPRESSED GENES AND GENE NETWORKS INVOLVED IN PIG OVARIAN FOLLICULAR ATRÉSIA

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1GenPhySE, Université de Toulouse, INRA, INPT, ENVT, Castanet Tolosan, France; 2INRA UMR 0085, CNRS UMR 7247, Université Francois Rabelais de Tours, IFCE, Physiologie de la Reproduction et des Comportements, Nouzilly, France; 3INRA, UMR1348 Pegase, Saint-Gilles, France; and 4AgroCampus-Ouest, UMR1348 Pegase, Saint-Gilles, France

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Terenina E, Fabre S, Bonnet A, Monniaux D, Robert-Granié C, SanCristobal M, Sarry J, Vignoles F, Gondret F, Monget P, Tosser-Klopp G. Differentially expressed genes and gene networks involved in pig ovarian follicular atresia. Physiol Genomics 49: 67–80, 2017. First published December 9, 2016; doi:10.1152/physigendis.00069.2016.—Ovarian folliculogenesis corresponds to the development of follicles leading to either ovulation or degeneration, this latter process being called atresia. Even if atresia involves apoptosis, its mechanism is not well understood. The objective of this study was to analyze global gene expression in pig granulosa cells of ovarian follicles during atresia. The transcriptome analysis was performed on a 9,216 cDNA microarray to identify gene networks and candidate genes involved in pig ovarian follicular atresia. We found 1,684 significantly regulated genes to be differentially regulated between small healthy follicles and small atretic follicles. Among them, 287 genes had a fold-change higher than two between the two follicle groups. Eleven genes (DKK3, GADD45A, CAMFA2, CCDC80, DAPK2, ECSIT, MSMB, NUPR1, RUNX2, SAMD4A, and ZNF628) having a fold-change higher than five could likely serve as markers of follicular atresia. Moreover, automatic confrontation of deregulated genes with literature data highlighted 93 genes as regulatory candidates of pig granulosa cell atresia. Among these genes known to be inhibitors of apoptosis, stimulators of apoptosis, or tumor suppressors INHBB, HNF4, CLU, different interleukins (IL5, IL24), TNF-associated receptor (TNFR1), and cytochrome-c oxidase (COX) were suggested as playing an important role in porcine atresia. The present study also enlists key upstream regulators in follicle atresia based on our results and on a literature review. The novel gene candidates and gene networks identified in the current study lead to a better understanding of the molecular regulation of ovarian follicular atresia. This process may be considered as physiologically normal, allowing the ovary to generate the ovulatory quota characteristic of the species, at each sexual cycle.

A number of studies in antral growing follicles have indicated that follicular atresia is initiated and caused by apoptosis (programmed cell death) of granulosa cells (53, 56, 71, 107). Two general pathways have been suggested to be involved in the physiological induction of apoptosis: negative induction by survival factor withdrawal and positive induction by the binding of a specific ligand, such as tumor necrosis factor-alpha or Fas ligand, to its membrane receptor (41). In follicles, apoptosis is triggered when endocrine and/or intra-follicular concentrations of survival factors, particularly gonadotropins and some growth factors, are inadequate (93). It is now recognized that a diverse spectrum of pro- and antiapoptosis susceptibility genes are thus expressed in germ cells and/or somatic cells of the ovary, including members of the BCL2 and caspase gene families (108). Many of these genes are regulated by gonadotropins and growth factors, and changes in the temporal pattern of cell death gene expression suggest that an intimate association exists between the products of these genes and the activation of apoptosis.

The biochemical characteristics of apoptosis have been investigated to discover causative factors and markers identifying follicles at the early stages of atresia. Results from studies of ovarian gene expression in vivo and in vitro have indicated that ovarian products, such as steroid hormones, transforming growth factor family members, insulin-like growth factor (IGF) 1, IGF-binding proteins (IGFBPs), and inhibins could play a role in atretic process (2, 42). However, most of this information has been acquired based on candidate-gene approaches.

In the present study, we propose a high-throughput transcriptomic approach to get a better understanding of the molecular actors, their regulatory genes, and physiological networks involved in pig follicle atresia. A specific microarray has been developed to identify genes expressed in granulosa cells along the terminal ovarian follicle growth in the pig (17). This represents a useful tool to study porcine ovarian follicular atresia. The aims of this study were 1) to analyze global gene expression changes in pig granulosa cells of ovarian follicles during atresia and 2) to identify the gene networks and propose new candidates genes that can...
further serve as targets to characterize and modulate atresia in granulosa cells.

**MATERIAL AND METHODS**

**Collection of ovaries.** Estrous cycles of 10 sows were synchronized by oral administration of 20 mg/day altenogest (Regumate; Hoechst-Roussel, Paris, France) for 18 days. Ovaries were removed by laparotomy 24 or 96 h after the last altenogest feeding. All procedures were approved by the Agricultural and Scientific Research agencies and conducted at an INRA experimental farm (animal experimentation authorization C37-175-2) in accordance with the guidelines for Care and Use of Agricultural Animals in Agricultural Research and Teaching.

**Granulosa cell isolation and RNA extraction.** Once ovaries were collected, all visible (>1 mm in diameter) antral follicles were isolated carefully under a binocular microscope. After dissection, follicle diameter was measured. Only follicles measuring 1–2 mm were kept, and granulosa cells were recovered from all individual follicles as described previously (78) and stored at −80°C until RNA extraction. For each follicle, a sample of granulosa cells was smeared on a histological slide and then stained by Feulgen reagent to determine follicular quality (76, 77). The quality of each follicle was assessed by microscopic examination of smears according to classical histological criteria; follicles were judged healthy (noted SHF for small healthy follicle), when having frequent mitosis and no pyknosis in granulosa cells, or atretic (noted SAF for small atretic follicle), when no mitosis and numerous pyknotic bodies in granulosa cells were observed. This method of classification has been previously demonstrated as particularly relevant since it very well reflects the level of expression of markers of follicular growth such as LH receptor, aromatase, and PAPP-A (73).

Within each sow, the follicles were pooled by quality classes, to generate SHF (small healthy follicle) and SAF (small atretic follicle) categories, respectively. Total RNA was extracted from SHF and SAF according to the technique described by Chomczynski and Sacchi (25) with minor modifications (46). The quality of each RNA sample was checked through the Bioanlyser Agilent 2100 (Agilent Technologies, Massy, France) and low-quality RNA preparations were discarded (RNA integrity number <3).

**Microarray hybridization.** After RNA quality check, 16 independent RNA samples were obtained, corresponding to 7 SHF samples and 9 SAF samples, respectively. A porcine microarray produced by CRB-GADIE (http://crb-gadie.inra.fr) and published in the Gene Expression Omnibus (GEO) database as GPL3729 (https://www.ncbi.nlm.nih.gov/projects/geo/query/acc.cgi?acc=GPL3729) was used to hybridize the RNA samples as previously described (17, 18). cDNA from luciferase was present on the array as positive control (193 spots), and water was included as negative control (64 spots). Microarrays were first hybridized with a 32P-labeled oligonucleotide sequence present in all PCR products to control the quality of the spotting process and to evaluate the amount of DNA in each spot (26). After being stripped, the arrays were hybridized with 32P-labeled complex probes synthesized from 5 µg of each RNA sample, using SuperScript II RNase H reverse transcriptase (Invitrogen, Cergy-Pontoise, France). Each complex probe was hybridized on membrane exposed for 1, 3, or 7 days at 68°C with a phosphor imaging system at 25 µm resolution (BAS-5000, Fujifilm, Raytest, Courbevoie, France). The imaging plates were scanned thereafter with a phosphor imaging system at 25 µm resolution (BAS-5000, Fujifilm, Raytest). Hybridization images obtained from oligonucleotide and complex probes were quantified using the semiautomated AGScan software (22, 26).

**Microarray data management.** The experimental design, its implementation, and the handling of data comply with MIAME standards (19), and all the experimental data were managed with BASE software (90), adapted by the Sigenae bioinformatics platform (http://www.sigenae.org) to manage radioactive experiments.

**Microarray data analysis.** Raw data and normalized data were submitted to the GEO database as GSM596969 (https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSM596969).

Data exploration and statistical analysis were performed using R software (http://www.r-project.org). The data coming from complex probes hybridization were analyzed on a natural logarithmic scale. As a first step, negative spots with >7 intensity values were checked on images and replaced by the median of negative spots in case of obvious overshining effect or hybridization stain. Then, luciferase-positive controls were removed, and the correlations of the gene expressions from each membrane were examined within condition. Hybridizations with a correlation coefficient <0.80 were discarded. Spots with low signal value (below the average of empty spots + 3 standard deviations) were considered as unexpressed and were excluded from the analysis. Finally, the negative control spots were removed and the remaining data were centered for each membrane.

The significance of the follicle status (SAF or SHF) on gene expression levels was assessed by a Student’s t-test (P < 0.05), followed by the Benjamini-Hochberg procedure controlling false discovery rate (FDR 5%) for each cDNA (9).

The relevance of the selection procedure was further evaluated via an unsupervised hierarchical clustering using the Ward method and Euclidian distance with the R functions hclust and heatmap (24).

**Sequence annotation.** Each cDNA sequence was compared with Refseq_rna mammalian database using the National Center for Biotechnology Information’s blastn program (http://blast.ncbi.nlm.nih.gov/blast.cgi). Blast results with an e-value <1e-3 were parsed and filtered to keep queries matching a gene or an mRNA or a CDS and possessing a global coverage of at least 70% of the query sequence. Resulting hits were sorted out according to their closeness to the pig genome, their coverage, and sequence identity. The selected cDNA sequences were submitted to the HUGO (Human Genome Organization) gene nomenclature committee, using their RefSeq IDs (http://www.genenames.org). Then, HUGO gene symbols were used as gene names.

**Quantitative PCR analysis of target gene expression.** Total RNA (1 µg) used in microarray experiments (SAF, n = 5; SHF, n = 5, among the samples used for array analysis) was reverse-transcribed as previously described (110). Primer sequences for genes were designed using Beacon Designer 7.0 software (Premier Biosoft Int., Palo Alto, CA) and are given in Table 1. The TCTP gene (transitionally controlled tumor protein, accession number BX667045) and MT-CO3 gene (cytochrome C oxidase subunit III, accession number CT971556) were used as internal controls (17). Quantitative real-time PCR was performed using SYBR green fluorescence detection during amplification on a Roche LC480 system (Roche Diagnostics, Meylan, France) according to the manufacturer’s recommendations. Duplicates of each template, consisting of 3 µl of a 1/10 dilution of the reverse transcription reaction, were loaded in 384-well plates with a 10 µl PCR mix SYBR green Power master mix (Applied Biosystems, Courtaboeuf, France), and 0.15 µM of forward and reverse primers (final volume 20 µl) were added. The PCR amplification conditions were as follows: 50°C for 30 min, initial denaturation at 95°C for 10 min, and 40 cycles (95°C for 15 s and 60°C for 1 min). The last cycle was followed by a dissociation step (ramping to 95°C). The real-time PCR amplification efficiency was calculated for each primer pair with five 1.2 dilution points of the calibrator sample (pool of the 10 cDNA samples) and was not lower than 1.8. After determination of the threshold cycle for each sample, the Pfaffl method was applied to calculate the relative changes of each mRNA in each sample (83). The relative expression was normalized by the corresponding geometric average of the two internal genes using geNorm v3.4 (113). The significance of differential expression between SHF and SAF was tested by Student’s t-test.

**Pathway analysis.** A pathway is an interconnected arrangement of processes, representing the functional roles of genes in the genome. The biological processes into individual genes may participate were.
1,684 cDNAs, 770 were upregulated in atretic follicles (SAF) groups corresponding to SAF and SHF status. Among the differentially expressed cDNAs separated the two experimental supervised hierarchical clustering (Fig. 1) shows that the 1,684 scripts corresponded to 1,223 unique HUGO genes. The unsupervised hierarchical clustering was established with respect to a null distribution constructed by permutations. Functional analysis was done using all genes that compose the array as “set reference.” A threshold network score of 40 as the highest score obtained with the help of the gene set reference was applied to select the highest significant networks for further analysis.

Search for upstream regulators. To propose upstream regulatory candidates that may have participated to differences in gene expression patterns between SAF and SHF, a recently developed web tool (http://keyregulatorfinder.genouest.org/) (13) was used. This methodology provides a reasonable number of upstream regulatory candidates from a submitted list of regulated genes, and success has been demonstrated for different case studies. In the present experiment, the differentially expressed genes were identified by their official gene symbol and submitted to analysis. The sign of variation was assigned to show the expression levels between SAF and SHF samples (+ + when upregulated in SAF vs. SHF, “−” when downregulated, and “??” when sign of variation between groups was not consistent for different probes of the same gene). The software output is a list of specific regulatory candidates. A posteriorization of the candidates was made by calculating the intersection with the differentially expressed genes in the input list, using Jvenn software (6).

RESULTS

Differentially expressed genes in follicle atresia. Follicles dissected from ovaries of cyclic sows were classified into two different categories based on individual diameter and quality traits assessed by histological examinations. Small healthy follicles (noted SHF) were then compared with small atretic follicles (noted SAF) for high throughput expression patterns. After normalization, 4,675 out of the 8,959 transcripts present on the pig cDNA array were found to be expressed in ovarian porcine follicles. Using the criterion of FDR < 5%, we identified 1,684 transcript spots as significantly regulated between SHF and SAF (Supplemental Table S1). (The online version of this article contains supplemental material.) These transcripts included 1,339 known cDNAs and 345 unknown cDNAs (Supplemental Table S1). The 1,339 annotated transcripts corresponded to 1,223 unique HUGO genes. The unsupervised hierarchical clustering (Fig. 1) shows that the 1,684 differentially expressed cDNAs separated the two experimental groups corresponding to SAF and SHF status. Among the 1,684 cDNAs, 770 were upregulated in atretic follicles (SAF) and 914 in healthy follicles (SHF). When we used a cut-off of |2| in fold-change between conditions, 142 genes were upregulated in SAF group (Table 2), whereas 145 genes were downregulated in those follicles compared with SHF (Table 3). In addition, some genes have undergone a very high fold-change (up to 17-fold) between follicle conditions, indicating that their expression level was very sensitive to atresia. About 11 genes notably exhibited a higher than fivefold overexpression in SAF and very poor expression in SHF (defined as belonging to the group of 20% of the lower expressed genes on the microarray) and were thus considered potential biomarkers of atresia in porcine small follicles: DKK3, GADD45A, CAMTA2, CCDC80, DAPK2, ECSIT, MSMB, NUPR1, RUNX2, SAMD4A, and ZNF628.

Quantitative PCR validation. Eight of these genes, chosen as being representative for differential expression levels in microarray between SHF and SAF, were further examined by quantitative (q)PCR (Fig. 2). As expected, higher expression levels for GADD45A, CLU, and HMOX1 were confirmed in SAF compared with SHF, whereas LRPR8, GSTA1, and CSNK2B exhibited decreased expression in the former compared with SHF. Notably, fold-change in expression levels for CLU, LRPR8, and GSTA1 between SHF and SAF categories was largely higher for qPCR than for microarray analysis (9.3 vs. 3.6, −26.9 vs. −2.7, and −42.3 vs. −10.5, respectively). The qPCR analysis also confirmed a trend to an increased expression level (P = 0.1) for CDKN1B in SAF compared with SHF. Finally, SH3GLB1, which was found as a nondifferentially expressed gene between experimental groups in microarray, did not differ between those groups in qPCR analysis. Thus, we confirm the reliability of the microarray approach for a set of genes with various expression ranges and profiles.

| Gene Name | Sense Primer (5’→3’) | Anti-sense Primer (5’→3’) |
|-----------|----------------------|--------------------------|
| CDKN1B    | ACGGGCTTTAGCCGAGGCCA | ACATACTTTAAATCCGAGCAGTTACG |
| CLU       | GCCAGGCTGAGAAGCTGATG | GGAATGACGGAGCGTTGTACCTCC |
| MT-Co3    | GCCAGGCTGAGAAGCTGATG | GCCAGGCTGAGAAGCTGATG |
| CSNK2B    | GGAGATTGAATTTGTCCTGGATGTAGTC | ACGCCCTGAAGTGAAGATGAG |
| GADD45A   | GGGTGGTCTGAGTGGTAGTATG | TGTGGCTGCTGCTGCTG |
| GSTA1     | CCGAGGAGATGAGGATGATC | TGGCGATGTAATTGAGGATGGC |
| HMOX1     | AGACCAGAAAGCGAGACGCC | ACCGGTTCCCAGGAAAGCC |
| LRPR8     | GCCCGGCAACAGATGCTGCAG | GCCCTCTTACAACACAGCTC |
| SH3GLB1   | GCCGCTGAGAAGAGAAGG | TTAGAGCTGTCTTGGAGGATGG |
| TCP1      | ATGAGCCTAGCCGAGCCATTTG | GCCGCTTGCTGCTGCTGTA |

TABLE 1. Sequences of oligonucleotide primers tested by quantitative RT-PCR analysis
death and survival, cell morphology, and reproductive system development and function and involves 61 genes (35 genes were upregulated and 26 genes were downregulated under atresia). Figure 3 shows nodes and edges of the first regulatory network related to endocrine system disorder, pointing on TP53 and N3RC1 genes as the central nodes.

Search for upstream regulators. The 287 differentially expressed genes (FDR < 5%, fold-change > 2|2| between SAF and SHF) were automatically confronted with the literature data on mammalian cell signaling to hierarchize upstream regulatory candidates. A dedicated web tool algorithm (http://keyregulatorfinder.genouest.org) was used, and a postprioritization of these candidates was made by comparing the list of candidates with the expression levels of these genes in SAH and SHF groups in the microarrays. Figure 4 shows the intersections of both lists; the detailed list of corresponding genes is in Supplemental Table S3. Among those regulatory candidates, 44 were included in the list of upregulated genes with fold-change $>$2 and 49 as downregulated genes with fold-change $<$2 between SAF and SHF conditions. Two genes, ESR1 (fold-change = $-1.9$) and NFKB1 (fold-change = $-1.3$), were present in the list of differentially expressed genes having a fold-change $<$2 between conditions. Nineteen genes were proposed as potential regulators, without being stated as differentially expressed by the microarray experiment. Among those 19 genes, only four genes (CSRP3, JUN, NEF2L2, and STAT1) were present on the pig GPL3729 microarray; however, their signal intensities were below the cut-offs; this does not preclude a regulatory role via posttranslational processes.

The remaining 15 genes (ADGRA3, ADGRE4P, ADGRG1, AP2A1, ATF2, BHLHE41, FOS, GST, IFRD1, IRF9, NFYA, RUNX1, SPI1, USF1, and USF2) were absent from the microarrays.

DISCUSSION

We used a comprehensive gene expression profiling by means of microarray analysis to identify groups of genes differently expressed by follicular status in pig ovaries. Similar microarray approaches have been used previously to characterize sheep granulosa cells and oocytes during early follicular development (16), growing healthy antral pig follicles (18), changes in granulosa cells gene expression associated with growth, plateau, and atretic phases in medium bovine follicles (29), and granulosa cells in the theca interna from bovine ovarian follicles during atresia (47–49). However, the present study is the first gene array analysis investigating ovarian follicular atresia in pigs. A large number of cDNA were found as differential between quality categories of follicles (healthy SHF vs. atretic SAF), and 287 differentially expressed annotated genes having a large fold-change between the two conditions were highlighted.

It has been previously proposed that apoptosis is the main biological process involved in follicular atresia (105, 106). This coordinated process involves many posttranslational processes, particularly the activation of a group of caspases. However, as illustrated by Fig. 5, a number of genes proposed by others as triggering apoptosis are not included in the list of
differentially expressed genes between SAF and SHF conditions in the present experiment, although they were present in the microarray design. It is important to note that previous studies did not necessarily evaluate the presence of those genes in a similar context. In comparison between bovine follicles in the plateau phase and atresia stage (29) classical granulosa apoptosis markers such as FAS (TNF receptor superfamily, member 6) and BCL2-associated X protein (BAX) are not significantly different in their expression. This result is supported by the findings of (47), showing that none of the caspases or Fas genes were differentially expressed in granulosa cells during atresia of bovine ovarian follicles. As previously shown (29), since follicle-stimulating hormone is thought to modulate the dynamics of the growth, plateau, and atretic phases in bovine follicles, this corresponds to the steady BAX levels studied phases. As reviewed in Hatzirodos et al. (47), most studies have been performed on antral follicles of a larger size (30, 54, 65) than those used in this experiment. It is possible that cell death mechanisms that operate in larger follicles are different from those at an earlier stage, where cells are under different hormonal control, LH in addition to FSH, and with focimatrix present (54). An alternative explanation is that only those granulosa cells in the atretic follicle with elevated expression of antiapoptotic genes are capable of surviving longer during the process of apoptosis (47, 88).

Table 2. List of upregulated genes under atresia

| HUGO Name | Fold-Change | Position |
|-----------|-------------|----------|
| MSMB      | 17.1        | 4246 SOX4 |
| NUPR1     | 14.6        | 1811 PTPRP |
| CAMTA2    | 7.5         | 6196 LRBP1 |
| ZNF628    | 7.2         | 5462 PLIN2 |
| RUNX2      | 6.9         | 3814 TMSB4 |
| ECSIT      | 6.2         | 5450 BHLHE49 |
| ARHGEF25  | 6.1         | 9097 LCN6 |
| GADD45A    | 5.5         | 6663 FTYL |
| SAMD4A     | 5.3         | 5745 EGR1 |
| CCDC80    | 5.2*        | 1752 VMP1 |
| DKK3       | 5.1         | 5863 PPP1R10 |
| DAPK2      | 5.0         | 5449 NUDFA1 |
| RASSF8     | 4.8         | 6052 MITF |
| SAFB2      | 4.5         | 3661 COMT |
| BCA51      | 4.4         | 3908 EPHA4 |
| EMR4P      | 4.0         | 3290 HMX01 |
| GPCR39     | 4.0         | 3146 CH3L1 |
| CD24       | 4.0         | 6225 NACC2 |
| RND3       | 4.0         | 2099 ISG15 |
| CAPG       | 3.8         | 4918 JUP |
| SPP1       | 3.8         | 5559 FLT1 |
| MMP2       | 3.7         | 6136 GPX3 |
| ZNF282     | 3.7         | 3897 ZNF395 |
| ZNF514     | 3.7         | 6208 LMF1 |
| CLU        | 3.6*        | 2586 ALCAM |
| CEBPD      | 3.5         | 8794 CDC152 |
| ZERI       | 3.5         | 7814 METTL7B |
| FBXO32     | 3.5         | 4996 THBS1 |
| FOSL2      | 3.5*        | 1184 ZNF609 |
| THAP1      | 3.4         | 4977 EFEMP2 |
| VAT1       | 3.4         | 771 CTNNA1 |
| VRK3       | 3.3         | 7801 EGF |
| IGFBBP3    | 3.3         | 2032 ITM2B |
| FAM134B    | 3.2*        | 2672 PLAT |
| PRRC2C     | 3.2         | 5758 PLEKHO1 |
| LGALS3     | 3.2         | 7363 PSMC2 |
|          |            |          |

| HUGO Name | Fold-Change | Position |
|-----------|-------------|----------|
|          |             |          |

Only fold-changes >2 between small atretic follicle (SAF) and small healthy follicle (SHF) categories are retained in this list. Position relative to the Gene Expression Omnibus (GEO) accession number GPL3729. *Means of several positions with the same annotation.

Functional networks involved in pig ovarian follicular atresia. In the present study, three networks of co-regulated differentially expressed genes were mainly involved in the molecular differences between SAF and SHF groups. These networks included both down- and upregulated genes, suggesting that cellular changes (notably related to “cancer” and “reproductive system disease”) may have occurred to counteract degenerative effects of other genes.

The first functional network is related to endocrine system disorders and reproductive system diseases. In this network, a decrease in the expression of some genes has been previously found to be associated with the atretic process. Particularly, we observed decreased expression levels of INHA, INHBA, and INHBB in SAF vs. SHF groups that is consistent with roles of inhibin family genes in atresia of granulosa cells. Inhibin inhibits pituitary secretion of follicle-stimulating hormone through negative feedback regulation (3, 14). Amounts of inhibin precursors increase initially in the largest follicles, but intrafollicular amounts of the large-molecular-weight forms do not change further with differentiation to dominance (3). Inhibins have a role in controlling follicular development, through either regulation of systemic gonadotropins or local intraovarian modulation of the effects of the gonadotropins (87). Previous research has shown that growing follicles contain increased total amounts of inhibin, whereas atretic and
Only fold-changes >2 between SAF and SHF categories are retained in this list. Ratio is inversed and preceded by a minus sign when value is <1. Position relative to the GEO accession number GPL3729. *Means of several positions with the same annotation.

subordinate follicles contain decreased amounts of the largest precursor forms. Further studies are required to elucidate this mechanism. Follicles destined to become atretic are characterized by loss of capacity to produce estradiol and enhanced production of low-molecular-weight IGFBPs. These changes not only precede atresia but also occur before the cessation of follicular growth and changes in intrafollicular inhibins (101).

Downregulation of these genes as well as the expression pattern was observed in various forms of atresia and programmed cell death (29). Bonnet et al. (18) previously showed overexpression of the genes implicated in lipid metabolism in granulosa cells during pig terminal follicular development. This finding is in agreement with differentiation mechanisms leading to fully steroidogenic granulosa cells in large antral follicles attested to by the overexpression of CYP19A1. This first functional network also includes upregulation of the transcription factor HNF4A (hepatocyte nuclear factor 4 alpha). In addition, there is a large increase in expression level of MSMB (beta-microseminoprotein) in SAF compared with the SHF group. This latter gene was rather downregulated in prostate carcinoma (27); downregulation of MSMB has been related to uncontrolled cell growth and enhance tumorigenesis (85, 102). Alterations in MSMB gene expression are associated with the development of prostate cancer (45). Upregulation of NUPR1 (nuclear protein 1) was also observed in SAF; this gene encodes proteins that inhibit apoptosis,
promotes pancreatic cancer development, and protects cells from stress (44), raising the question of its role in atretic follicles. A role for AKT2, coding for kinases controlling cell proliferation mechanism, has been also previously observed in cancers (7, 118). The ADAMTS1 gene family is related to the cleavage of extramembranous domains (92). The expression of this gene is increased significantly during follicular growth and atresia in small but not large follicles, and more in preovulatory follicles of older than younger cows (57).

The second network is related to cancer and to cell death and survival. Many of these genes participate to the control of the actin microfilament network, including myosin-related genes. This may reflect changes in cell shape that generally accompany the atresia process in follicles. Actin is associated with

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Fig. 2. Real-time quantitative (q)PCR validation of expression levels of target genes found differentially expressed between SHF and SAF by microarray analyses. Total RNA from pig granulosa cells of small healthy follicles (SHF, \( n = 5 \)) and small atretic follicles (SAF, \( n = 6 \)) were reverse-transcribed and submitted to real-time qPCR analysis for quantification of SH3GLB1, CLU, GADD45A, HMOX1, CDKNB1, CSNK2B, LRP8, and GSTA1 gene expression levels. Quantitative qPCR data (black histogram) are means ± SE of relative expression to the reference genes TCTP and MT-CO3. PCR data are compared with normalized microarray data (white histogram) where SH3GLB1 was chosen as a nondifferentially expressed gene. Significant difference in qPCR analysis between means from SHF and SAF were obtained after a Student’s t-test (after Welch correction for unequal variance): \( * P < 0.05, *** P < 0.001 \).
granulosa cell shape changes during antral folliculogenesis (18) that may consequently affect steroid synthesis and proliferation. This network also includes two upregulated genes in the SAF category, \textit{CEBPD} and \textit{CLU}, which are involved in cell death, tumor progression, and neurodegenerative disorders (55, 120). Among others, the growth arrest and DNA damage-inducible, alpha (\textit{GADD45A}) gene has been recently reported as an indicator of late atresia (29). Indeed, \textit{GADD45A} activity is required for the progression of apoptotic cell death in follicular atresia, as it controls cell cycle arrest, apoptosis induction, and DNA damage repair in response to DNA-damaging agents (119). It has also been shown that transfection of a \textit{GADD45A} expression vector has induced apoptosis, reportedly by interacting MEKK4/MTK1 and activating JNK/p38 signaling, which induces apoptosis (103). Douville and Sirard (29) have suggested that \textit{GADD45A} activity is increasingly needed for the progression of apoptotic cell death in follicular atresia. These results highlight the fact that a particular gene expression pattern per stage reflects follicle growth or atresia.

In addition, the \textit{HIF1A} gene was found to be downregulated in atretic follicles. It encodes the alpha subunit of transcription factor hypoxia-inducible factor-1 and plays an essential role in embryonic vascularization, tumor angiogenesis, and pathophysiology of ischemic disease. \textit{HIF1A} induction is mediated by \textit{IGF1} in cells grown under normal oxygen concentrations, thus uncoupling \textit{HIF1A} induction from oxygen deprivation (34). Proapoptotic and antitumorigenic effects of \textit{HIF1A} have also been reported (4). Finally, mutations in \textit{RUNX2} (run-related transcription factor 2), a gene that was upregulated in SAF vs. SHF categories, have been correlated with cleidocranial dysplasia (40) and associated with a prostate cancer (84). Upregulation in \textit{DKK3} induces apoptosis in cisplatin-resistant lung adenocarcinoma cells via the Wnt/\beta-catenin pathway (115). Another important gene in this network may be \textit{DAPK2}, a member of the death-associated protein kinase (DAPK) family and related to apoptosis pathways (8). This gene was also reported to be a modulator of TRAIL [tumor necrosis factor (TNF)-related apoptosis-inducing ligand] signaling (5). Therefore, upregulated \textit{DAPK2} gene in atretic follicles suggests a DAPK2-mediated role of apoptosis process (36) in atresia.

The third network concerns cell death and survival, cell morphology, and reproductive system development and function. Among these genes, two are particularly interesting. The ECSIT signaling integrator was identified as a mitochondrial protein that has a neuroapoptotic role in Alzheimer’s disease pathogenesis (99). \textit{CCDC80} (coiled-coil domain containing 80) plays a role in cell organization and apoptosis, and inhibition of \textit{CCDC80} expression may notably be an important event in the development of colorectal and pancreatic cancers (15, 39).

Our results are in concordance with our previous study where we described gene sets whose expression increases during the follicular development and decreases with atresia. The atresia process requires fewer new genes than the growth process (98). Moreover, our finding corresponds to differentially expressed genes along the terminal follicular growth described by Bonnet and colleagues (18). This study showed in particular downregulation of ribosomal protein, cell morphology, and ion-binding genes. Numerous genes whose expression increased during the terminal follicular growth

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**Fig. 3. Atresia biomarker-transcription factor connections.**

**Fig. 4. Graphic representation of the various gene lists.** Differentially expressed genes [false discovery rate (FDR) 5%], either upregulated genes [fold-change (FC) $>2$], or downregulated between SHF and SAF groups (FC $< -2$), and specific regulatory upstream candidates as suggested by automatic confrontation with literature data (KeyRegulatorFinder web tool) are colored in green, blue, red, and yellow respectively. The number of common genes is indicated for any of the intersection between these 4 lists.
Fig. 5. Apoptosis signaling pathway that highlights the key molecular events involved in triggering apoptosis. Genes in gray correspond to the genes present on the used array but not showing as differentially expressed in our study.
were implicated in glutathione metabolism (GSTA1) and lipid metabolism (CYP19A). One of predictors is actin, which is associated with granulosa cell shape changes along antral folliculogenesis that may consequently affect steroid synthesis and proliferation.

Altogether, these genes could form the basis for a future study to develop a panel of tissue indicators with the purpose of prognosis of this phenomenon.

Regulators and markers of porcine follicle atresia. If the genes are coexpressed during follicular atresia, it is highly probable that they are co-regulated, so a better knowledge of upstream actors responsible for the regulation of gene coexpression modules may provide new insights into molecular mechanisms controlling atresia. A study of upstream transcriptional regulators may provide additional information that can explain the observed gene expression changes in the list of studied genes. Such an investigation is made arduous by the fact that the key upstream regulator may be not included in the list of genes suggested as differentially expressed according to a threshold probability (13). This approach was successfully used in our previous study to propose upstream transcriptional regulators that may participate in molecular flexibility in porcine adipose tissues in response to diet (38). A large number of differentially expressed genes in our study encoded transcription factors. Moreover, using academic knowledge on reactions and regulations included in TRANSPATH database (59) and a dedicated algorithm to search and sort upstream regulatory molecular actors (13), we were able to propose upstream regulators to different genes and biological processes. Figure 4 summarizes the differentially expressed genes including upregulated genes (fold-change > 2) and downregulated genes (fold-change < -2) between noted SHF and SAF, together with potential specific candidates acting as upstream regulators of these genes. This analysis notably highlights 93 genes as possible regulatory candidates of pig granulosa cell atresia, for which expression changes larger than twofold between conditions were also experimentally observed between SHF and SAF groups. Whereas most of them have not yet been described in follicular atresia, they are generally known as inhibitors of apoptosis, stimulators of apoptosis, or tumor suppressors. Among them, INHBB, HNF4, and CLU were again in evidence. In addition, different interleukins (IL5, IL24), TNF-associated receptor (TNFRI), and cytochrome-c oxidase (COX) may have played a key role in porcine atresia. Table 4 summarizes key

Table 4. Key upstream regulators in follicle atresia

| HUGO Name | Description | Reference No. |
|-----------|-------------|---------------|
| IMPDH2    | inosine 5'-phosphate dehydrogenase 2 | (61) |
| GATA4     | GATA binding protein 4 | (60) |
| MEF2      | myocyte enhancer factor 2 | (1) |
| GSTA1     | glutathione S-transferase alpha 1 | (16, 31, 89, 111) |
| GADD45A   | growth arrest and DNA-damage-inducible, alpha | (50, 119) |
| BOC       | cell adhesion associated, oncogene regulated | (63) |
| CTSI      | cathepsin L | (116) |
| NF-YA     | nuclear factor Y-box A | (20, 35, 80) |
| IL5       | interleukin 5 | (43, 68, 69, 72) |
| IL24      | interleukin 6 | |
| IL6       | interleukin 24 | |
| IL6R      | interleukin 24 | |
| Nfkb1     | nuclear factor of kappa light polypeptide gene enhancer in B-cells 1 | (10) |
| COX       | cytochrome C | (52, 71, 96) |
| ID1       | inhibitor of DNA binding 1 | (91) |
| INHBB     | inhibin beta B | (64, 112) |
| CD47      | transmembrane glycoprotein | (67) |
| NDUF56    | NADH dehydrogenase (ubiquinone) Fe-S protein 6 | |
| PPAR isotypes | peroxisome proliferator-activated receptors | (121) |
| RPN2      | ribophorin II | (33) |
| SORT1     | sortilin 1 | (37) |
| TNFR1     | TNF receptor 1 | (58) |
| CALM1     | calmodulin 1 | (100) |
| MET1      | metaxin 1 | (21) |
| SDHB      | succinate dehydrogenase complex, subunit B | (23) |
| TSPO      | translocator protein | (86) |
| HNF4a     | hepatocyte nuclear factor 4, alpha | (62) |
| FDP5      | farnesyl diphosphate synthase | (66) |
| CLU       | clusterin | (12, 55, 70, 79, 104, 117, 120) |
| CEBP      | CCAAT/enhancer binding protein | |
| MAP1A, MAP1B | microtubule-associated protein | (32) |
| HINT1     | histidine triad nucleotide binding protein 1 | (11) |
| HNF1A     | HNF1 homeobox A | (51) |
| ZNF350    | zinc finger protein | (28) |
| GADD45A   | growth arrest and DNA-damage-inducible, alpha | (81, 94, 109) |
| CD82      | transmembrane glycoprotein | |
| SOX4      | SRY (sex determining region Y)-box 4 | (122) |
| ARF       | ADP-ribosylation factor | (97, 114) |
upstream regulators in follicle atresia based on our results and on a literature review.

Besides the upstream regulatory actors of pig follicle atresia, the present study also highlights 11 genes (DKK3, GADD45A, CAMTA2, CCDC80, DAPK2, ECSIT, MSMB, NUPR1, RUNX2, SAMD4A, ZNF628) with very sensitive expression to atresia (fold-change >5 in atretic vs. healthy follicles). All these genes belong to networks identified above and are linked to molecular functions related to proliferation of tumor cell lines and transcription regulation. It is thus proposed that they could further serve in a panel of tissue prognosis indicators of porcine follicle atresia. Six genes (GADD45A, RUNX2, CCDC80, DKK3, NUPR1, DAPK2) among these 11 genes and described above are functionally connected by two other transcription factors (NR3C1 and TP53) as illustrated in Fig. 3. The NR3C1 (nuclear receptor subfamily 3, group C, member 1) gene encodes a glucocorticoid receptor, which can function both as a transcription factor that binds to glucocorticoid response elements in the promoters of glucocorticoid-responsive genes and as a regulator of other transcription factors. NR3C1 expression has been previously demonstrated to have importance in cancer (82). As previously described (47), we suggest an association between atretic status and genes that are influenced by the p53 transcription factor. TP53 encodes the tumor protein p53, which responds to diverse cellular stresses to regulate target genes inducing cell cycle arrest, activation of apoptosis, senescence, DNA repair, or changes in metabolism (74). The tumor suppressor protein p53 has a critical role in regulation of senescence, DNA repair, or changes in metabolism (74). The present study also enlists key upstream regulators in follicle atresia based on our results and on a literature review. Identification of 93 upstream regulatory candidate genes, which are known to regulate genes encoding key regulators of cell cycle, apoptosis, and proliferation in follicle atresia. These markers could further serve in a panel of tissue prognosis indicators of porcine follicle atresia.

AUTHOR CONTRIBUTIONS

GTK, PM, SF, AB, ET and DM discussed the aims of the study and the data retrieval. AB, JS, FV contributed to the data collection and data retrieval. MSC and CRG developed and performed all the statistics. FG performed the upstream regulators study. ET, GTK, PM, SF, AB, MSC, CRG and DM contributed to the interpretation and discussion of the results. ET drafted the manuscript and all authors contributed to this manuscript in its final version. All authors read and approved the final manuscript.

REFERENCES

1. Akhtar MW, Kim MS, Adachi M, Morris MJ, Qi X, Richardson JA, Bassel-Duby R, Olson EN, Kavalali ET, Montegia LM. In vivo analysis of MEF2 transcription factors in synapse regulation and neuronal survival. PLoS One 7: e34863, 2012. doi: 10.1371/journal.pone.0034863.
2. Ashkenazi A, Disit VM. Death receptors: signaling and modulation. Science 281: 1305–1308, 1998. doi: 10.1126/science.281.5381.1305.
3. Austin EJ, Mihm M, Evans AC, Knight PG, Ireland JL, Ireland JJ, Roche JF. Alterations in intrafollicular regulatory factors and apoptosis during selection of follicles in the first follicular wave of the bovine estrus cycle. Biol Reprod 64: 839–848, 2001. doi: 10.1095/biolreprod.64.3.839.
4. Bacon AL, Harris AL. Hypoxia-inducible factors and hypoxic cell death in tumour physiology. Annu Med 36: 530–539, 2004. doi: 10.1080/07853904010018231.
5. Bai T, Tanaka T, Yukiowa K. Targeted knockdown of death-associated protein kinase gene expression induces TRAIL-mediated apoptosis in human endometrial adenocarcinoma cells. Int J Onco 37: 203–210, 2010. doi: 10.3892/ijo.0000668.
6. Bardou P, Mariette J, Escudié F, Djemiel C, Kloczko C. jven: an interactive Venn diagram viewer. BMC Bioinformatics 15: 293, 2014. doi: 10.1186/1471-2105-15-293.
7. Bellacosa A, de Feo D, Godwin AK, Bell DW, Cheng JQ, Altomare DA, Wang M, Dubeau L, Scambia G, Masciullo V, Ferrandina G, Benedetti Panici P, Mancuso S, Nerl G, Testa JR. Molecular alterations of the AKT2 oncogene in ovarian and breast carcinomas. Int J Cancer 64: 280–285, 1995. doi: 10.1002/ijc.2910640412.
8. Benderska N, Schneider-Stock R. Transcription control of DAPK. Apoptosis 19: 298–305, 2014. doi: 10.1007/s10495-013-0931-6.
9. Benjamini Y, Hochberg Y. Controlling the False Discovery Rate: A Practical and Powerful Approach to Multiple Testing. J R Stat Soc B 57: 289–300, 1995.
10. Bernal GM, Wahlstrom JS, Crawley CD, Cahill KE, Pytel P, Liang H, Kang S, Weichselbaum RR, Yamini B. Loss of Nfkb1 leads to early onset aging. Aging (Albany NY) 6: 931–943, 2014. doi: 10.18632/aging.100702.
11. Beyoğlu D, Krausz KW, Martin J, Maurhofer O, Dorow J, Ceglarek U, Gonzalez FJ, Dufour JF, Idle JR. Disruption of tumor suppressor gene Hint1 leads to remodeling of the lipid metabolic phenotype of mouse liver. J Lipid Res 55: 2309–2319, 2014. doi: 10.1194/jlr.M050682.
12. Birkenmeier EH, Gwynn B, Howard S, Jerry J, Gordon JI, Landschulz WH, McKnight SL. Tissue-specific expression, developmental regulation, and genetic mapping of the gene encoding CCAAT/enhancer binding protein. Genes Dev 3: 1146–1156, 1989. doi: 10.1101/gad.3.8.1146.
13. Blazy P, Gondret F, Lagarrigue S, van Milgen J, Siegel A. Using a large-scale knowledge database on reactions and regulations to propose key upstream regulators of various sets of molecules participating in cell metabolism. BMC Syst Biol 8: 32, 2014. doi: 10.1186/1752-0509-8-32.
14. Bleach EC, Glencross RG, Feist SA, Groome NP, Knight PG. Plasma inhibin A in heifers: relationship with follicle dynamics, gonadotropins, and steroids during the estrous cycle and after treatment with bovine follicular fluid. Biol Reprod 64: 743–751, 2001. doi: 10.1095/biolreprod64.3.743.
15. Bommer GT, Jäger C, Dürr EM, Baehs S, Eichhorst ST, Brabletz T, U, Gonzalez FJ, Dufour JF, Idle JR. Disruption of tumor suppressor gene Hint1 leads to remodeling of the lipid metabolic phenotype of mouse liver. J Lipid Res 55: 2309–2319, 2014. doi: 10.1194/jlr.M050682.
16. Bonnet A, Blevinsca C, Benne F, Bodin L, Cotinot C, Lianbet L, Sancristobal M, Serr J, Terenina E, Martin P, Rosler-Klopp G, Mandon-Pepin B. Transcriptome profiling of sheep granulosa cells and oocytes during early follicular development obtained by laser capture
yis of pancreatic cancer reveals a tumor suppressor function for HNF1A. Carcinogenesis 35: 2670–2678, 2014. doi:10.1093/carcin/bgu193.

52. Hsu SY, Hsueh AJ. Tissue-specific Bel-2 protein partners in apoptosis: An ovarian paradigm. Physiol Rev 80: 593–614, 2000.

53. Hughes FM Jr, Gorospe WC. Biochemical identification of apoptosis (programmed cell death) in granulosa cells: evidence for a potential mechanism underlying follicular atresia. Endocrinology 129: 2415–2422, 1991. doi:10.1210/endo-129-5-2415.

54. Irving-Rodgers HF, Harland ML, Rodgers RJ. A novel basal lamina matrix of the stratified epithelium of the ovarian follicle. Matrix Biol 23: 207–217, 2004. doi:10.1016/j.matbio.2004.05.008.

55. Jin W, Li Q, Wang J, Chang G, Lin Y, Li H, Wang L, Gao W, Pang T. Na+/H+ exchanger 1 inhibition contributes to K562 leukemic cell differentiation. Cell Biol Int 36: 739–745, 2012. doi:10.1002/cbin.2010091.

56. Jolly PD, Smith PR, Heath DA, Hudson NL, Lun S, Still LA, Watts CH, McNatty KP. Morphological evidence of apoptosis and the prevalence of apoptotic versus mitotic cells in the membrana granulosa of ovarian follicles during spontaneous and induced atresia in ewes. Biol Reprod 56: 837–846, 1997. doi:10.1093/biolreprod56.4.837.

57. Khan DR, Fournier É, Dufoi I, Richard FJ, Singh J, Sirard MA. Meta-analysis of gene expression profiles in granulosa cells during folliculogenesis. Reproduction 151: R103–R110, 2016. doi:10.1530/REP-15-0594.

58. Kim IV, Song EH, Lee HJ, Oh YK, Choi KH, Yu DY, Park SI, Seong HJ, Park JA, Moreno CS. Global transcriptome analysis of formalin-fixed prostate cancer specimens identifies biomarkers of disease recurrence. PLoS One 8: e709 –717, 2010. doi:10.1371/journal.pone.0010382.

59. Kyrölähti A, Kauppinen M, Lind E, Unkila-Kallio L, Butzow R, Kim JY, Song EH, Lee HJ, Kato H, Hashidume T, Shimizu M, Sato R. Biochemical identification of apoptosis (programmed cell death) in granulosa cells: evidence for a potential new role in ovarian follicle development. Endocrinology 142: 5243–5253, 2001. doi:10.1210/endo.142.12.8517.

60. Meek DW. Regulation of the p53 response and its relationship to cancer. Biochem J 469: 325–346, 2015. doi:10.1042/BJ20150517.

61. Miyashita T, Krajewska S, Krajewska M, Wang HG, Lin HK, Liebermann DA, Hoffman B, Reed JC. Tumor suppressor p53 is a regulator of bc1 and bax gene expression in vitro and in vivo. Oncogene 9: 1799–1805, 1994.

62. Monpetet M, Moniaux D, Monniaux D. Short-term effects of FSH in vitro on granulosa cells of individual sheep follicles. J Reprod Fertil 79: 505–515, 1987. doi:10.1530/jf.0.0790505.

63. Moniaux D, Pisselet C. Control of proliferation and differentiation of ovine granulosa cells by insulin-like growth factor-I (IGF-I), IGF-II, and their binding proteins during growth and atresia of ovine ovarian follicles. Endocrinology 132: 1438–1446, 1993. doi:10.1210/endo.132.4.7681760.

64. Monniaux D, Pisselet C. Control of proliferation and differentiation of ovine granulosa cells by insulin-like growth factor-I and -like-stimulating hormone in vitro. Biol Reprod 46: 109–119, 1992. doi:10.1095/biolreprod46.1.109.

65. Nakagawa H, Liyanarachchi S, Davuluri RV, Auer H, Martin EW Jr, de la Chapelle A, Frankel WL. Role of cancer-associated stromal fibroblasts in metastatic colon cancer to the liver and their expression profiles. Oncogene 23: 7366–7377, 2004. doi:10.1038/sj.onc.1208013.

66. Norquay LD, Lin Y, Surabhi RM, Gietz RD, Tanese N, Cattini PA. A member of the nuclear factor-1 family is involved in the pituitary regulation of the human placental growth hormone genes. Biochem J 354: 387–395, 2001. doi:10.1042/BJ2000387.

67. Ono M, Handa K, Withers DA, Hakomori S. Motility inhibition and apoptosis are induced by metastasis-suppressing gene product CD82 and its analogue CD82, with concurrent glycosylation. Cancer Res 59: 2335–2339, 1999.

68. Patel AS, Karagas MR, Perry AE, Spencer SK, Nelson HH. The role of short-term effects of FSH in vivo on granulosa cells of individual sheep follicles. J Reprod Fertil 79: 505–515, 1987. doi:10.1530/jf.0.0790505.

69. Pfeiffer MW. A new mathematical model for relative quantification in real-time RT-PCR. Nucleic Acids Res 29: e45, 2001. doi:10.1093/nar/29.9.e45.

70. Pi M, Quarels LD. GPRC6A regulates prostate cancer progression. Prostate 72: 399–409, 2012. doi:10.1002/pros.21442.

71. Pomerantz MM, Shrestha Y, Flavin RJ, Regan MM, Penney KL, Mucci LA, Stuempfle MJ, Hunter DJ, Chakravorty S, Saha R, Wedge DJ, Olsen JL, Tabernero J, Baselga J, Richardson AL, Loda M, Oh WK, Kantoff PW, Hahn WC, Freedman ML. Analysis of the 10q11 cancer risk locus implicates MSMB and NCOA4 in human prostate tumorigenesis. Proc Natl Acad Sci 110: 8866–8871, 2013. doi:10.1073/pnas.1307978110.

72. Roche JF. Control and regulation of folliculogenesis—a symposium in perspective. Rev Reprod 19: 1–27, 1996. doi:10.1530/rr.0.0010009.

73. Rodgers RJ, Lavranos TC, van Wezel II, Irving-Rodgers HF. Development of the ovarian follicular epithelium. Mol Cell Endocrinol 151: 171–179, 1999. doi:10.1016/S0303-7207(99)00087-8.

74. Roméro I, Andrews K, Pouvreau K, Menon A, Kirby G. Human GSTA1-1 reduces c-Jun N-terminal kinase signalling and apoptosis in Caco-2 cells. Biochem J 400: 135–141, 2006. doi:10.1042/BJ20060110.
104. Sutcliffe S, De Marzo AM, Sfanos KS, Laurence M.

TARGET GENES OF PIG OVARIAN FOLLICULAR ATRESIA

91. Sayasith K, Lussier J, Siros J. Molecular characterization and transcriptional regulation of a disintegrin and metalloproteinase with thrombospondin motif 1 (ADAMTS1) in bovine preovulatory follicles. Endocrinology 154: 2857–2869, 2013. doi: 10.1210/en.2013-1140.

95. Schenfeld N, Bauer MK, Grimm S.

90. Sarre LH, Troein C, Vallon-Christersson J, Gruvberger S, Borg A, Petersson C. BioArray Software Environment (BASE): a platform for comprehensive management and analysis of microarray data. Genome Biol 3: SOFTWARE0003, 2002. doi: 10.1186/gb-2002-3-8-software0003.

91. Sato AY, Antonioli E, Tambellini R, Campos AH. ID1 inhibits USP2 and blocks TGF-β-induced apoptosis in mesangial cells. Am J Physiol Renal Physiol 301: F1260–F1269, 2011. doi: 10.1152/ajprenal.00128.2011.

92. Sayasith K, Lussier J, Siros J. Molecular characterization and transcriptional regulation of a disintegrin and metalloproteinase with thrombospondin motif 1 (ADAMTS1) in bovine preovulatory follicles. Endocrinology 154: 2857–2869, 2013. doi: 10.1210/en.2013-1140.

93. Scaramuzzi RJ, Baird DT, Campbell BK, Driancourt MA, Dupont J, Fortune JE, Gilchrist RB, Martin GB, McNatty KP, McNelly AS, Monget P, Monniaux D, Văinescu D, Webb R. Regulation of folliculogenesis and the determination of ovulation rate in ruminants. Reprod Fertil Dev 23: 444–467, 2011. doi: 10.1071/RD09161.

94. Schoenfeld N, Bauer MK, Grimm S. The metastasis suppressor gene C33/C2D2/KAI1 induces apoptosis through reactive oxygen intermediates. FASEB J 18: 159–160, 2004. doi: 10.1096/fj.03-04200fp.

95. Schuler M, Green DR. Mechanisms of p53-dependent apoptosis. Biochem Soc Trans 28: 689–688, 2001. doi: 10.1042/bst0290684.

96. Schultz DR, Harrington WJ Jr. Apoptosis: programmed cell death at a molecular level. Semin Arthritis Rheum 32: 345–369, 2003. doi: 10.1053/sarh.2003.50005.

97. Sherr CJ. Divorcing ARF and p53: an unsettled case. Nat Rev Cancer 6: 663–673, 2006. doi: 10.1038/nrc1954.

98. Sirard MA, Toward building the cow folliculome. Anim Reprod Sci 149: 90–97, 2014. doi: 10.1016/j.anireprosci.2014.06.025.

99. Soler-López M, Badiola N, Zanzoni A, Aloy P. Towards Alzheimers root cause: ECSIT as an integrating hub between oxidative stress, inflammation and mitochondrial dysfunction. Hypothetical role of the adapter protein ECSIT in familial and sporadic Alzheimers disease pathogenesis. BioEssays 34: 532–541, 2012. doi: 10.1002/bies.201100193.

100. Stanislaus A, Bakhtiar A, Salleh D, Tiasah S, Fatemian T, Hossain S, Fernandes D, Salleh D, Tiash S, Fatemian T, Hossain S, Amin A, Peterson C. BioArray Software Environment (BASE): a platform for comprehensive management and analysis of microarray data. Genome Biol 3: SOFTWARE0003, 2002. doi: 10.1186/gb-2002-3-8-software0003.

101. Tilly JL, Tilly KI, Perez GI. The genes of cell death and cellular susceptibility to apoptosis in the ovary: a hypothesis. Cell Death Differ 4: 180–187, 1997. doi: 10.1038/sj.jdd.4400238.

102. Tolley H, Barrett JC. CD82 metastasis suppressor gene: a potential target for new therapeutics? Trends Mol Med 11: 563–570, 2005. doi: 10.1016/j.molmed.2005.10.002.

103. Tosser-Klopp G, Bonnet A, Yerle M, Hatey F. Functional study and regional mapping of 44 hormono-regulated genes isolated from a porcine granulosa cell library. Genet Sel Evol 33: 69–87, 2001. doi: 10.1186/1297-9686-33-1-69.

104. Tosser-Klopp G, Benne F, Bonnet A, Mulsant P, Gasser F, Hatey F. A first catalog of genes involved in pig ovarian follicular differentiation. Mamm Genome 8: 250–254, 1997. doi: 10.1007/s003359900403.

105. Tu F, Pan ZX, Yao Y, Liu HL, Liu SR, Xie Z, Li QF. mir-34a targets the inhibin beta B gene, promoting granulosa cell apoptosis in the porcine ovary. Genet Mol Res 13: 2504–2512, 2014. doi: 10.4238/2014.January.14.6.

106. Vandesompele J, De Preter K, Pattyn F, Poppe B, Van Roy N, De Paepe A, Speleman F. Accurate normalization of real-time quantitative RT-PCR data by geometric averaging of multiple internal control genes. Genome Biol 3: RESEARCH0034, 2002. doi: 10.1023/b:mirb.00000277-research0034.

107. Wartmann T, Mayerle J, Kähne T, Sahin-Tóth M, Ruthenbürger M, Lippert H, Schulz HU, Volanakis EJ, Sherr CJ. Developmental strategies for evasion of Arf tumor suppression. Cell Cycle 9: 14–15, 2010. doi: 10.1016/j.ccr.2009.11.0390.

108. Wang Z, Ma L, Kang Y, Li X, Zhang X. Dickkopf-3 (Dkk3) induces apoptosis in cisplatin-resistant lung adenocarcinoma cells via the Wnt/β-catenin pathway. Oncol Rep 33: 1097–1106, 2015. doi: 10.3892/or.2014.3704.

109. Wang Z, Ma L, Kang Y, Li X, Zhang X. Dickkopf-3 (Dkk3) induces apoptosis in cisplatin-resistant lung adenocarcinoma cells via the Wnt/β-catenin pathway. Oncol Rep 33: 1097–1106, 2015. doi: 10.3892/or.2014.3704.

110. Wartmann T, Mayerle J, Kähne T, Sahin-Tóth M, Ruthenbürger M, Lippert H, Schulz HU, Volanakis EJ, Sherr CJ. Developmental strategies for evasion of Arf tumor suppression. Cell Cycle 9: 14–15, 2010. doi: 10.1016/j.ccr.2009.11.0390.

111. Wartmann T, Mayerle J, Kähne T, Sahin-Tóth M, Ruthenbürger M, Lippert H, Schulz HU, Volanakis EJ, Sherr CJ. Developmental strategies for evasion of Arf tumor suppression. Cell Cycle 9: 14–15, 2010. doi: 10.1016/j.ccr.2009.11.0390.

112. Wang Z, Ma L, Kang Y, Li X, Zhang X. Dickkopf-3 (Dkk3) induces apoptosis in cisplatin-resistant lung adenocarcinoma cells via the Wnt/β-catenin pathway. Oncol Rep 33: 1097–1106, 2015. doi: 10.3892/or.2014.3704.

113. Wang Z, Ma L, Kang Y, Li X, Zhang X. Dickkopf-3 (Dkk3) induces apoptosis in cisplatin-resistant lung adenocarcinoma cells via the Wnt/β-catenin pathway. Oncol Rep 33: 1097–1106, 2015. doi: 10.3892/or.2014.3704.

114. Wang Z, Ma L, Kang Y, Li X, Zhang X. Dickkopf-3 (Dkk3) induces apoptosis in cisplatin-resistant lung adenocarcinoma cells via the Wnt/β-catenin pathway. Oncol Rep 33: 1097–1106, 2015. doi: 10.3892/or.2014.3704.