miRNA-Profiling in Ejaculated and Epididymal Pig Spermatozoa and Their Relation to Fertility after Artificial Insemination

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Simple Summary: The present study searched for the presence and abundance of porcine spermatozoa small RNA sequences (microRNAs) that have the potential to alter gene expression patterns. Four different sperm sources were compared: spermatozoa from three different sections of the ejaculate and from the caudal epididymis, also classed as spermatozoa from higher (HF) or lower (LF) fertility boars. Sperm miRNAs were compared using high-output small RNA sequencing. We identified five sperm miRNAs not previously reported in pigs. Differences in abundance of four miRNAs known to affect the expression of genes with key roles in fertility were related to boar fertility. These miRNAs could be used as fertility markers in artificial insemination programs.

Abstract: MicroRNAs (miRNAs) are short non-coding RNAs (20–25 nucleotides in length) capable of regulating gene expression by binding -fully or partially- to the 3’-UTR of target messenger RNA (mRNA). To date, several studies have investigated the role of sperm miRNAs in spermatogenesis and their remaining presence toward fertilization and early embryo development. However, little is known about the miRNA cargo in the different sperm sources and their possible implications in boar fertility. Here, we characterized the differential abundance of miRNAs in spermatozoa from the terminal segment of the epididymis and three different fractions of the pig ejaculate (sperm-peak, sperm-rich, and post-sperm rich) comparing breeding boars with higher (HF) and lower (LF) fertility after artificial insemination (AI) using high-output small RNA sequencing. We identified five sperm miRNAs that, to our knowledge, have not been previously reported in pigs (mir-10390, mir-10386, mir-10390, mir-6516, mir-9788-1, and mir-9788-2). Additionally, four miRNAs (mir-1285, mir-92a, mir-34c, mir-30), were differentially expressed among spermatozoa sourced from ejaculate fractions and the cauda epididymis, and also different abundance was found between HF and LF groups in mir-182, mir-1285, mir-191, and mir-96. These miRNAs target genes with key roles in fertility, sperm survival, immune tolerance, or cell cycle regulation, among others. Linking the current findings with the expression of specific sperm proteins would help predict fertility in future AI-sires.

Keywords: spermatozoa; miRNAs; fertility biomarkers; epididymis; ejaculate; pig

1. Introduction

The expression of more than 60% of all genes is estimated to be under the regulation of non-coding small RNAs (ncsRNAs) [1]. Currently, several families of ncsRNAs: microRNAs (miRNAs), short interfering RNAs (siRNAs), piwi-interacting RNAs (piRNAs), and transfer RNA (tRNA) have been identified within somatic cells [2–4], with the miRNA
family being the best described among the species studied so far. The miRNAs are endogenous ncsRNA molecules of 20–25 nucleotides (nt) in length, capable of altering the translation of their messenger RNA (mRNA) targets [5], to influence protein production [6–8] by regulating protein translation during the cell cycle [9]. The abundance of several miRNAs has been described in mammalian spermatozoa (both in epididymal and ejaculated spermatozoa [10]). Specifically, studies of miRNAs in spermatozoa searched for their role in spermatogenesis and their remaining presence toward fertilization and early embryo development, attempting to disclose if their presence or absence in mature spermatozoa might be related to abnormal development and functioning [11,12] and/or fertility modulation, as found in the murine species where the transfer of sperm miRNAs into the oocyte after fertilization was associated with differential early embryo development [3,13–16]. Moreover, Dicer1 (an enzyme required for miRNA synthesis) in deficient mice developed morphologically abnormal spermatids, low sperm motility, and low fertility [17]. In the porcine genome, up to 457 mature miRNAs have been identified and annotated to date (miRBase 22, www.mirbase.org (accessed 2 March 2018)), among which, several of these miRNAs have been previously studied for expression and functionality [18,19]. However, much remains to be unveiled regarding the identification and biological functions of miRNAs in pig spermatozoa, [12,20], and of differences in the expression of specific miRNAs among semen samples, with different sperm morphology and/or motility. Furthermore, pig spermatozoa recovered from the epididymis terminal section (EpiTS, e.g., mature spermatozoa in the cauda and the convoluted first portion of the vas deferens) [21] or from the ejaculate, depicted miRNAs with the ability to influence the expression of target genes partially responsible for spermatogenesis, sperm maturation, and zygote development [22,23]. The boar ejaculate consists of three different major fractions: the pre-sperm fraction (pre-SRF, with absence of spermatozoa), the sperm–rich fraction (SRF), and the post-sperm-rich fraction (post-SRF). The SRF can be further divided into a sperm-peak fraction (SPF), consisting of the first 10–15 mL of the SRF holding approximately 25% of the total ejaculate sperm numbers, and being rich in epididymal caudal fluid [24], and the remaining SRF, which maintains an elevated sperm concentration and protein secretions from the vesicular and prostate glands. The post-SRF not only presents relatively fewer spermatozoa but also increasing secretions of vesicular, prostate, and bulbourethral glands [25]. In the present study, we report deep sequencing of miRNAs from spermatozoa collected from three fractions of the ejaculate (ejaculated spermatozoa exposed to different secretions of the accessory sexual glands, namely SPF, rest of SRF and post-SRF) of mature breeding boars. Furthermore, since miRNAs have been suggested as potential biomarkers in breeding programs [26], the boars used in this study were classified as having higher (HF) or lesser (LF) fertility subsequent to commercial artificial insemination (AI) as well as spermatozoa retrieved from their EpiTS. Our aim was the genome-wide identification and profiling of miRNAs in the different sperm sources to disclose which miRNAs were present after sperm maturation and whether they changed after ejaculation. Moreover, since miRNAs are crucial for fertilization and preimplantation embryo development [27,28], we further postulate that although many factors can influence male fertility, the identification of specific miRNAs could help its prognosis. Therefore, we additionally analyzed whether a differential abundance of specific miRNAs between boars with higher- or lower-fertility could have a putative role as fertility biomarkers, by their relation to potential gene targets. Selected preliminary data had been reported elsewhere [29].

2. Materials and Methods

2.1. Ethics Statement

All experiments were accomplished in accordance with the European Directive 2010/63/EU EEC for animal experiments and accepted by the University of Murcia’s Re-
search Ethics Commission (research code: 639/2012) and the “Regional Committee for Ethical Approval of Animal Experiments” (Linköpings Djurförsöksstiska nämnd) in Linköping, Sweden (permits Dnr 75-12, ID1400 and 03416-2020).

2.2. Experimental Design

All boars used in this study (n = 6) presented normal semen quality. The miRNA expression profile was studied in in ejaculated spermatozoa manually collected from each of three ejaculate fractions (SPF, rest of SRF, and post-SRF) as well as in epididymal spermatozoa collected from the EpiTS from the same boars, slaughtered due to genetic renewal at the enterprise. The ejaculates (n = 18) were monthly collected (one collection/boar/month) in within 3 months. In this same 3-month period, other ejaculates were weekly collected as routine in the AI center and used to prepare commercial insemination doses for farms. In total, 923 sows were inseminated, and after farrowing, both farrowing rate (FR, proportion of inseminated sows that farrowed) and litter size (LS, number of piglets born per farrowing) were recorded. The raw AI-fertility data, provided by Topics Norsvin España (Madrid, Spain), were subjected to statistically correction for factors associated to farm and sow to separate the individual contribution of each boar to fertility; accounted as the deviation of FR (in percentage) and LS (in number) of each boar [29,30]. The boars were then, according to their FR (high-fertility boars: FR > 0.94; low-fertility boars: FR < −0.42) or LS (high-fertility boars: LS > 0.11; low-fertility boars: LS < −0.14), classified as having higher (HF; n = 3, 540 AIs) or lower (LF; n = 3, 315 AIs) fertility. The miRNAs from HF and LF boar spermatozoa (ejaculated and epididymal) were compared.

2.3. Boars Handling and Sample Collection

Spermatozoa from ejaculates and the epididymis terminal section (EpiTS) used in the experiment were collected from six healthy, mature cross-bred boars (Landrace × Large-White breeds; 2–3 years old) of known fertility, housed in a commercial AI-station (Topigs Norsvin España, Calasparra, Murcia, Spain). To ensure good semen production over time, the boars were housed in a building with constant lighting 16 h per day and equipped with evaporative coolers to maintain temperature and air humidity within comfortable ranges.

Using the gloved-hand method, three separate fractions of the ejaculate (SPF, SRF and post-SRF) were collected from each boar over a period of three months (one ejaculation per boar and month). All ejaculates used in the experiment fulfilled the requirements of sperm quantity and quality to produce commercial AI-doses (> 200 × 106 spermatozoa/mL, >70% progressive motile and >75% with normal morphology). Considering the ejaculates were fractioned, the concentration of spermatozoa was measured (SP-100 NucleoCounter; ChemoMetec A/S, Allerød, Denmark) in a recomposed sample (mixing specific aliquots of the fractions) while sperm motility was assessed using CASA system (ISASVi® CASA; Proiser R + D, Valencia, Spain) and sperm morphology assessed with phase contrast microscopy. The results were compared to data of previous and posterior ejaculates, and they were found to be non-deviating. Each of the three ejaculate fractions was double centrifuged (1500×g at room temperature (RT) for 10 min (Rotofix 32A; Hettich Centrifuge UK, Newport Pagnell, Buckinghamshire, England, UK), to harvest the spermatozoa. The boars were slaughtered after being culled for genetic improvement decisions while they were still healthy and fertile, to collect the contents of their EpiTS via cannulation [30]. All sperm pellets obtained were kept in 15 or 2 mL-tubes at −80 °C until further analyses.

2.4. RNA Extraction

Total RNA was isolated from spermatozoa using a miRNeasy kit (Qiagen, Hilden, Germany) designed to isolate RNA for low quantity samples following the manufacturer’s instructions. Briefly, each sample (200 µL) was thawed on ice and incubated with 200 µL
of QIAzol Lysis Reagent (Qiagen, Hilden, Germany) at room temperature (RT) for 2 h. Samples were mixed by pipetting and vortexing, treated with 100 μL of chloroform, hand-shaken (RT, 15 s) and incubated at RT for 3 min. Then, samples were centrifuged (12,000× g, 4 °C, 15 min) and the aqueous phase (200 μL) was mixed with 300 μL of 100% ethanol. Samples were placed into a RNeasy MinElute spin column (Qiagen, Hilden, Germany) in a 2 mL collection tube and centrifuged (8000× g, RT, 15 s). Several washes were performed to the column (500 μL of RWT buffer (8000× g, RT, 15 s), 500 μL of RPE buffer (8000× g, RT, 15 s), and 500 μL of 80% ethanol (8000× g, RT, 2 min), discarding the flow-through after each washing. Subsequently, the column was centrifuged (15,000× g, RT, 5 min) and the flow-through was discarded. The RNA was eluted by centrifuging the spin column membrane containing 10 μL nuclease-free water (15,000× g, RT, 1 min). Total RNA content and its quality was determined by NanoDrop® 1000 (Thermo Fisher Scientific, Waltham, MA, USA). RNA concentration in the samples ranged from 50 to 100 ng/μL and an A260/A280 ratio of ~1.8 was achieved in all samples, indicating good RNA purity. Samples were kept at ~80 °C until further analysis.

2.5. Small RNA Library Preparation

Small RNA libraries were built using the Illumina TruSeq Small RNA Sample kit (RS-200-0012 and RS-200-0024: Indexes 1-24, Illumina, San Diego, CA, USA) following the manufacturer’s instructions. The aim of this procedure was to ligate adapters to each end of the RNA molecule, and then reverse-transcribe and amplify to generate a cDNA library. A gel purification step prepared the library for clustering and sequencing. Briefly, 1 μg total RNA in 5 μL of molecular grade water was used for each sample. The 3' and 5' adapters were ligated to the samples by incubation at 28 °C for 1 h in a thermal cycler. Then, samples were subjected to reverse transcription (50 °C for 1 h) followed by amplification by PCR to create cDNA constructs based on the small RNA ligated with 3' and 5' adapters. This step selectively enriched RNA fragments with adapter molecules on both ends. Amplification was performed with 2 primers that annealed to the adapter ends. The PCR settings for the amplification step were as follows: 98 °C for 30 s, 14 cycles of 98 °C for 10 s, 60 °C for 30 s, 72 °C for 15 s, and then 72 °C for 10 min and held at 4 °C. Each library was then run on a High Sensitivity DNA chip (Agilent Technologies, Santa Clara, CA, USA). A total of 24 libraries were then pooled in equal molar amounts and run in a 6% TBE gel (Life Technologies, Carlsbad, CA, USA) at 140 V for 55 min. Bands between 140 and 160 bp containing miRNAs were excised from the gels. These gel pieces were centrifuged in a gel breaker tube (IST Engineering, Milpitas, CA, USA) at 20,000× g for 2 min to move the gel through the holes of the gel breaker tube into the 2 mL tube and incubated on a rotating chamber at RT for 2 h in 200 μL molecular grade water. The cDNA construct was then checked for quantity and quality with the Agilent High Sensitivity DNA Kit (Agilent Technologies, Santa Clara, CA, USA). For clustering, the total of all molarities from the peaks observed in the Bioanalyzer were used and the libraries were normalized to 2 nM using Tris-HCl 10 mM, pH 8.5. The cDNA construct was then denatured and clustered on a single read Illumina V2 flow cell (Illumina, San Diego, CA, USA) and ran on the Illumina NextSeq sequencing platform (NextSeq 500/550; Illumina, San Diego, CA, USA) with a Mid Output kit v2.5, with at least 10 million reads per sample for 150 cycles for with a 6-cycle indexing read.

2.6. Overview of Sequencing Performance

A total of 24 small-RNA libraries were performed (Boars n = 6 × Sperm sources n = 4) (six boars and four sperm sources per boar), which were internally quality filtered using the procedure chastity pass filter (PF, i.e., the ratio of the brightest base intensity divided by the sum of the brightest and second brightest base intensities) (Illumina System Guide (15050091 v03)) that Illumina’s NextSeq sequencer performs to calculate the percent of
passing on patterned and non-patterned Illumina flow cells. The NextSeq flow cell contained four physical lanes and the pooled library was loaded in all lanes. The outcomes of the sequencing process are depicted in Table 1 and Supplementary File S1.

Table 1. Sequencing outcomes for RNA-Seq procedure.

|               | Total Reads (n) | Pf Reads (%) | Reads >q30 (%) | Undetermined Reads (%) |
|---------------|-----------------|--------------|----------------|------------------------|
| Average Lanes 1–4 | 14,494,752      | 97.37        | 98.39          | 2.62                   |

2.7. Bioinformatic Analyses

Prior to alignment, reads were trimmed for 3’ and 5’ adapters using Trimmomatic (version 0.36) [31]. The ENCODE miRNA-Seq data were processed using STAR aligner v. 2.4.2a [32]. Clean reads were aligned on porcine genome (ssc10.2), and miRNA-Seq data were studied using Partek Genomics Suite 7.0 (Partek). A Principal Component Analysis (PCA) was performed on all samples (Supplementary File S1). Data were first normalized using the total count normalization method [33–36]. Differential expression of miRNAs among sperm sources was established by using a one-way ANOVA, setting parameters as a fold change (FC) > 1 or < −1 with p-value < 0.05. The raw datasets were deposited at Sequence Read Archive (SRA) with the BioProject accession number: PRJNA762225 (https://www.ncbi.nlm.nih.gov/sra/PRJNA762225 accessed on 12 March 2018).

2.8. Target Gene Prediction and Functional Analysis

In the present study, miRDB (http://mirdb.org (accessed on 12 March 2018)) was used to predict which potential miRNA targets could the differentially expressed sperm miRNAs relate to. For target prediction analyses, the miRNAs were aligned with their human homologous considering that the miRNA-target database has been established exclusively for human and some model organisms. Only target genes with a score > 90 (miRBase) were selected.

The network of biological functions and pathways based on the GO and KEGG databases was investigated using Cytoscape Software v3.0.0 (http://www.cytoscape.org/ (accessed on 12 March 2018)) application ClueGO v2.0.3.

3. Results

3.1. Identification of miRNAs in Spermatozoa from Cauda Epididymis and Ejaculate Fractions

Table 2 lists the miRNAs identified in boar spermatozoa that showed differences in abundance among sperm sources, highlighting six miRNAs that have not been described before in pig spermatozoa and five miRNAs that were significantly dysregulated (p < 0.005).

Table 2. List of identified miRNAs with different abundance among spermatozoa from three ejaculate fractions: sperm peak fraction (SPF), sperm rich fraction (SRF), and post-SRF of mature boars (n = 6), and the epididymal terminal segment (EpiTS).

| Group | miRNAs (n) | Name of miRNAs                  |
|-------|------------|---------------------------------|
| SPF   | 27         | mir-182, mir-6516, mir-29a, mir-10386, mir-let-7f-1, mir-30c-2, mir-30b, mir-96, mir-148a, mir-92a-1, mir-92a-2, mir-30d, mir-10390, mir-9788-1, mir-9788-2, mir-1285, mir-34c-1, mir-30e, mir-30c-1, mir-10a, mir-10391, mir-1839, mir-10b, mir-664, mir-132, mir-6782, mir-16-1 |
| SRF   | 26         | mir-9831, mir-148a, mir-96, mir-6516, mir-30e, mir-15b, mir-21, mir-221, mir-182, mir-1285, mir-100, mir-9788-1, mir-9788-2, mir-92a-1, mir-34c-1, mir-191, mir-... |
miRNAs marked in bold represent not previously described miRNAs in porcine spermatozoa. miRNAs marked in red represent miRNAs with significantly (p < 0.05) dysregulated abundance.

Figure 1 depicts the miRNAs commonly expressed among spermatozoa recovered from the three ejaculate fractions studied: SPF (n = 27), SRF (n = 26), and the post-SRF (n = 18), and from the EpiTS (n = 27), and where all sources shared the expression of 13 miRNAs.

### 3.2. miRNAs Were Differentially Expressed among Spermatozoa Ejaculate Fractions and from Cauda Epididymis, as Well as between Boars with Higher (HF) or Lower (LF) Fertility: Assessment of Chromosome Location and Structure

Some miRNA abundance differed significantly (p < 0.05) among sperm sources (Figure 2). The mir-1285 was upregulated in spermatozoa from the EpiTS compared to spermatozoa derived from any of the ejaculate fractions studied, as well as when comparing the SPF and the post-SRF fractions. The SPF-spermatozoa showed a higher expression of mir-92a-1, mir-92a-2, and mir-34c than spermatozoa from the SRF. The mir-30e was upregulated in SRF-spermatozoa compared with post-SRF spermatozoa (Table 3). Additionally, there was an overexpression of mir-191 and mir-96 (in spermatozoa from EpiTS and SRF, respectively), while a repression of mir-182 and mir-1285 was observed in spermatozoa from the SPF and the SRF, respectively, when comparing spermatozoa from boars of higher- or lesser fertility (Table 3). Figure 2 depicts chromosome location and precursor
structure of those miRNAs that were differentially expressed. Figure 3 depicts a hierarchical clustering of the pattern followed by all miRNAs in spermatozoa from different sources within all boars (n = 6) and in boars with higher- or lower-fertility (n = 3) within the same sperm source.

Table 3. Differentially abundant microRNAs (miRNAs, p-value < 0.05) in spermatozoa collected from the three ejaculate fractions studied (SPF: sperm-peak fraction; SRF: sperm-rich fraction; and post-SRF) and from the functional epididymal terminal segment (EpiTS) among all boars or comparing high-and low fertility boars.

| Group             | Name of miRNAs Differentially Abundant | Fold Change |
|-------------------|---------------------------------------|-------------|
| SPF vs. SRF       | mir-34c, mir-92a-1, mir-92a-2          | 2.5, 4, 4.1 |
| SRF vs. Post-SRF  | mir-30e                                | −1.2        |
| EpiTS vs. SPF     | mir-1285                               | 1.7         |
| EpiTS vs. SRF     | mir-1285                               | 1.6         |
| EpiTS vs. Post-SRF| mir-1285                               | 1.8         |
| HF vs. LF SPF     | mir-182                                | 1.1         |
| HF vs. LF SRF     | mir-96                                 | −3.3        |
| HF vs. LF Post-SRF| mir-1285                               | 1.1         |
| HF vs. LF EpiTS   | mir-191                                | −3.1        |

Figure 2. Chromosome location (left image) and miRNA precursor structure (right image) of all differentially expressed microRNAs (miRNAs) (p-value < 0.05 and ≥1.0 Fold Change (FC) or ≤−1.0) presented in this study: mir-182 (A,a), mir-191 (B,b), mir-34c-1 (C,c), mir-96 (D,d), mir-92a (E,e), mir-30e (F,f), mir-1285 (G,g). The images were created in Rfam version 14.5 [37]. The black color in the miRNA structure represents a template model, while differences between the template and the sequences are highlighted in color, depending on whether it is a modification (green), an insertion (pink), or a reposition (blue). Structures were generated by R2DT using the d.5.e.P.waltl template provided by CRW [38], copyright © 2022 Rfam Team.
3.3. Target Prediction and Functional Annotations of Differentially Expressed Sperm miRNAs

To gain insight into the biological function of the differentially expressed sperm miRNAs found in this study, we identified which target genes whose expressions could be potentially and/or partially regulated by these sperm miRNAs. We focused on those target genes with a score ≥90 of homology (Tables 4 and 5) in the miRNA-target human
database (miRDB), against which the miRNAs were aligned owing to the highly conserved degree of homology among species. A total number of 246 potential target genes likely influenced by the expression of the five miRNAs (miR-1285, miR-92a-1, miR-92a-2, miR-34c, and miR-30e) were found after comparing epididymal spermatozoa with those from ejaculate fractions (Table 4). Additionally, a total of 208 target genes were listed as putatively regulated by four miRNAs (miR-191, miR-182, miR-96, and miR-1285) found comparing spermatozoa from higher- with lower-fertile boars (Table 5). The networks representing interactions between those GO terms and biological pathways are shown in Figure 4. ClueGO software (Cytoscape) revealed the considered miRNA-targeted genes (target genes scoring > 90 in miRbase) in several immune-related pathways and cellular processes.

Table 4. List of predicted target genes of differentially expressed miRNAs (p value < 0.05 and > 1.0-Fold Change (FC) or < −1.0-FC) in spermatozoa from three different ejaculate fractions: sperm peak fraction (SPF), sperm rich fraction (SRF), or post-SRF, or from the epididymal terminal segment (EpiTS).

| Comparison | miRNA | FC | Predicted Targets (n) | Name of Predicted Targets |
|------------|-------|----|-----------------------|---------------------------|
| SPF vs. SRF | miR-92a-1 | 4-14 | SLX4 SH3TC2 LMLN NRP2 BCAM PRKCA ATN1 TOB1 ARM7 ANKS1A SFC3A ATP2B2 SLC2A1 COL1A1 | IQSEC2 FOXP4 SYNGAP1 BCL2A1 FAM22A NFIX TOMIL2 HNF4A NKAIN1 DLK1 STX1B MIER2 ZBTB4 NACC1 APH1A SLC6A17 WNT11 AR CELF5 PPP2R5B RAMP2 KCNQ4 BCAM TGM2 FIBCD1 CELSR2 SLC4A1 NOVA2 ZNF574 PDE1B DNAJB5 ZNF385A ZSWIM4 TF3 LRC4 L1CAM MTA1 CLIP2 PPPI9RB GIPC1 EHMT2 B4GALT1 COL5A3 TSC1 MLLT6 CAMTA1 DUSP3 FAM155B PBX2 SKI SMG5 RIM53 CAMK2A PAPP2 C120R43 NFC SLC22A11 HYOU1 SLA2 TNR6A DAMP SQCD GNAS NFAS NT5E | |
| SPF vs. Post-SRF | miR-34c | 2.5-85 | SORL1 AASDH ADGRG2 PDE4D PDCD6IP SGC8 ANKRD17 SCP2 TWIST2 SNAP25 ADF DAZAP1 ZNF483 C8orf58 GPBP1 L1 ZNF454 PA5 RX7 | HCN3 FAM76A MDM DLL1 FBKP1B SYT1 E2F5 PPP1R11 RAP1GDS1 FAM167A SDK2 SATB2 SCN2B MYCN NECST1 CELF3 MGAT4A LGR4 NAV3 NAV1 MET FLOT2 XYL1T1 AHCYL2 TGIF2 PCAS1 PKP4 CACNA1E MLT3 FUT9 RRAS PTP1CN1 MMP2 VAMP2 ABR SLC2A17 ATPX1 | |
| SRF vs. Post-SRF | miR-30e | -1.2-0 | SORL1 AASDH ADGRG2 PDE4D PDCD6IP SGC8 ANKRD17 SCP2 TWIST2 SNAP25 ADF DAZAP1 ZNF483 C8orf58 GPBP1 L1 ZNF454 PA5 RX7 | ACTR3C ADAM19 ADAMTS3 ADAMTS9 ADRA2A ALG10 ANKHD1 ANKR2A ANKRD17 ANO4 ASB3 ATG12 AZIN1 B3GNT5 BD1P BRD1 BRWD3 C9orf72 CALCR CARF CCDC61 CCDC43 CCDC97 CCNE2 CNT2 CELSR3 CFL2 CHD1 CHIC1 CHL1 CHST2 CLOCK CNKSR2 CNOT9 COL1A3 COL25A1 CYPA241A DCTN4 DCUN1D3 DDAH1 DES2 DIF5 DOLP1 E2F7 EEA1 EED ELL1 EML1 EML4 EXT2 FAM160B1 FAP BFOX45 FBKP3 FND3C AFOX1 FRMPD1 FRZB FZD3 GABRB1 GATN7 GMNC GOLGA1 HCFEC2 HDAC9 ITGA6 ITPK1 KIAA0408 KL10 KL12 KLH12 LCLAT1 LHX8 LIMCH1 LIN28B LMBR1 LMBR1L LPGAT1 LLRC17 MARC6 MAST4 MEJ0B MEOX2 MIER3 MKRN3 MLXIP MTDH MYH11 NAV3 NCA1 MECAP1 NEDD4 NEURL1B NFATS NFIB NTSE OTUD6B PAPOLA PCDH7 PDE7A PEX5 PFN2 PHPT2 PIP4K2A PLAGL2 PLEKH3 |
| Comparison          | miRNA   | Predicted Targets (n) | Name of Predicted Targets                                                                 |
|---------------------|---------|-----------------------|-------------------------------------------------------------------------------------------|
| EpiTS vs. Post-SRF  | miR-1285| 1.8                   | SORL1 AASDH ADGRG2 PDE4D PDCD6IP SGC8 ANKRD17 SCP2 TWIST2 SNAP25 AFP DAZAP1 ZNF483 C8ORF58 GPBP1L1 ZNF454 PAX5 RFX7 |
| EpiTS vs. SRF       | miR-1285| 1.6                   | SORL1 AASDH ADGRG2 PDE4D PDCD6IP SGC8 ANKRD17 SCP2 TWIST2 SNAP25 AFP DAZAP1 ZNF483 C8ORF58 GPBP1L1 ZNF454 PAX5 RFX7 |
| EpiTS vs. LF-SRF    | miR-1285| 1.8                   | SORL1 AASDH ADGRG2 PDE4D PDCD6IP SGC8 ANKRD17 SCP2 TWIST2 SNAP25 AFP DAZAP1 ZNF483 C8ORF58 GPBP1L1 ZNF454 PAX5 RFX7 |
| EpiTS vs. LF-SPF    | miR-182 | 98                    | PRKACB RGS17 BNC2 SNX30 LPP MITF FRS2 CAMSAP2 HAS2 PRRG3 EPAS1 PALLD DCUN1D1 SLC39A9 VAM3 PMP3 MTSS1 NPTX1 NEXMIF CD2AP TECTB PRDG SPATA13 SPIN1 CACNB4 MFAP3 CTTN NCAUD ACTR2 ABHD13 ADCY6 FOXF2 CADM2 TAF4B EDNRB RAB10 RAPgef5 LRCH2 CHIC1 ZF36L1 PCNX1 MAST4 ITPR1 RASA1 LMTK2 USP13 FOXO3 ZC3H15 MAGI1 TAGLN3 RARG LHX1 GNAQ LMS1 GXYLT1 NR1I2 STK19 IGFR1 CBF2T3 FLOT1 HOX9 BRF3 BSL5 FAM171A1 MEDI MYRIP TRABDB2 PYGO2 PPM1L KIAA1217 HOOK3 S1C2 BCL2L12 G1T2 BRMS1L PHP1 TMEM47 MGA1 NDFC3B BNIP3 ZFCH31 INT5S DCUN1D3 SLC35G1 PURA PPI1L1 SERTAD4 EV5 EV5 ADD3 L1CAM BMT2 STAG1 PLPPR4 ADGRL2 YWHAG HDAC9 ZNF483 ZNF483 REV1 |
| HF- vs. LF-SRF      | miR-96  | 73                    | ZCCH3 CUBN1 PALLD FOX2 CHST1 MYRIP ZTB41 FRMD5 CACNA2D2 PRRG3 SH3KBP1 NOV2 E2B1 MTOR SLCA9A1 PRGG3 TTYH3 NGL2 FOXO1 ARHGA6 ANKRD27 SENN3 CEP170B VAT1L PPP4R3A STA1A1 CD164 UNC13C DOCK1 SPEN TMEM170B REV1 PPM1L NR1N1 MGA1 STK19 TMEM198 SAST |
| HF- vs. LF-SPF      | miR-1285| 18                   | SORL1 AASDH ADGRG2 PDE4D PDCD6IP SGC8 ANKRD17 SCP2 TWIST2 SNAP25 AFP DAZAP1 ZNF483 C8ORF58 GPBP1L1 ZNF454 PAX5 RFX7 |
| HF- vs. LF-EpiTS    | miR-191 | 4                     | NEURL4 TAF5 CREBB CASTOR2                                                                 |

*Only target genes with a score >90 (miRbase) were selected.

**Table 5.** List of predicted target genes of differentially expressed miRNAs (*p* value < 0.05 and >1.0-fold change (FC) or <−1.0-FC) in spermatozoa from ejaculate fractions (spem-peak fraction (SPF), sperm-rich fraction (SRF), and post-SPF), and the functional epididymal terminal segment (EpiTS), and of higher fertility (HF) compared to lower fertility boars (LF).
4. Discussion

Multiple lines of evidence indicate that sperm miRNAs have crucial gene-regulatory roles in many fundamental biological processes, including spermatogenesis, prompting...
for sperm motility, fertilization, and early embryo development [14,15,19]. Although pig spermatozoa have been shown to express miRNAs ruling sperm motility, structural integrity and metabolism before [12,39], the present study aimed to perform comprehensive miRNA expression profiling of pig spermatozoa retrieved from different sources using a genome-wide deep sequencing approach. The results have provided novel findings regarding the miRNA content and its differential abundance in the spermatozoa from the EpiTS and from three ejaculate fractions (SPF, SRF, and post-SRF), for their plausible implication in sperm-related biological processes post-fertilization, ultimately influencing fertility by controlling the expression of specific genes.

After sperm production in the testis, spermatozoa acquire motility and fertilizing capacity during their journey along the epididymis [29], interacting with different ncsRNAs (including miRNAs), transcripts, and proteins that are released from the epithelium of the epididymis mostly via epididymosomes [40–42]. We have hereby identified a total of 27 miRNAs in spermatozoa from the EpiTS and the SPF, 26 in the SRF, and only 18 in the post-SRF. Interestingly, the differences between the numbers of commonly expressed miRNAs was not large between the SPF and the SRF (n = 19) or between EpiTS and SRF (n = 16), which seems reasonable considering the SRF (including its SPF) presents higher contents of epidydimal fluid [43].

Some of the miRNAs found in this study have not been described in pig spermatozoa before (mir-10386, mir-10390, mir-10391, mir-9788-1, mir-9788-2, and mir-6516, present in all sperm sources). mir-10386, mir-10390 and mir-10391 have only been described in porcine liver [44], mir-9788-1 and mir-9788-2 were identified in sow milk exosomes [45], while mir-6516 has been reported in other tissues in several species (porcine liver [44], human platelets [46], chicken embryos, [47] or mouse brain [48]. Further experimental evidence is needed to validate the presence and to identify the roles ascribed to these miRNAs in pig spermatozoa.

Few of the miRNAs identified in the present study were differentially expressed among sperm sources. Mir-1285 was found upregulated in spermatozoa from the epididymis terminal segment compared to spermatozoa from all of the different ejaculate fractions.

This miRNA was previously identified by our group in spermatozoa retrieved from the SRF [49], but, to the best of our knowledge, the present study is the first to report an overexpression of this miRNA in spermatozoa from the EpiTS compared to spermatozoa from the ejaculate fractions. The fact that mir-1285 abundance differed when compared to the other sperm sources could be explained by the fact that, during spermatogenesis, post-transcriptional control of gene expression is highly active [50] and mir-1285 appears to be involved in spermiogenesis by inhibiting boar Sertoli cell proliferation through the regulation of AMPK [51]. Phosphorylated AMPK is implicated in the regulation of 17β-estradiol-mediated inhibition of Sertoli cell viability through increasing p53 and p27 expression and inhibiting mTOR and Skp2 expression [51]. Moreover, mir-1285 regulates the expression of many genes associated with reproductive processes, for instance, the potential target gene DAZAP1 seems essential for normal development of spermatozoa in mice [52]. This finding supports the theory that miRNAs may play key roles in spermatogenesis. Additionally, we observed an overexpression of three miRNAs (mir-92a-1, mir-92a-2, and mir-34c) in SPF compared with the SRF spermatozoa. The differences in miRNAs abundance between sperm from different parts of the ejaculate suggests that miRNAs regulate mRNA expression in the final processing of spermatozoa during ejaculation [22]. MiR-92 has been shown to play a role in early chicken gonadogenesis by regulating the expression of the ATRX and DDX3X genes. The mir-92a target genes are implicated in the regulation of the NOTCH signaling pathway, which regulates many cellular events: inhibition of cell differentiation, cell proliferation, and preservation of stem cell population [53]. In addition, NOTCH genes are considered potential targets for mir-34 [54]. The mir-34c belongs to a family of evolutionarily conserved miRNAs (mir-34a, mir-34b, and mir-34c) with active influence on genes that control the cell cycle [55,56], including cell cycle
arrest, cellular senescence [57], and apoptosis [58]. Furthermore, the abundance of mir-34c is regulated by the p53 signaling pathway and could constitute a central inhibitor of p53 functions, which are responsible of a variety of intrinsic and extrinsic stress signals that impact upon cellular homeostatic mechanisms regulating DNA replication, chromosome segregation, and cell division [58]. mir-34c has been suggested as fundamental for the development of both male and female bovine gametes and therefore suggested as a potential biomarker of male bovine fertility [59]. Moreover, decreased levels of mir-34c have been observed in samples of infertile individuals [60]. Among the mir-34c target genes, three genes downregulated by mir-34c (TGF2, E2F5, and BMP3), all implicated in the regulation of Transforming Growth Factor-beta (TGF-β) signaling pathway drew our attention.

A recent study pointed the TGF-β signaling as a crucial mechanism for immune tolerance to spermatozoa, a very relevant capacity present in the epididymis to tolerate spermatozoa in the lumen, despite they present xenoantigens, which could cause a response from the male immune system [61]. After sperm deposition in the female reproductive tract, similar concepts would apply for the TGF-β signaling pathway, considering its role in preventing spermatozoa from autoimmune responses [49], while simultaneously, the female immune system provides effective protection against ascending pathogens [62]. The genes mentioned above are transcriptional repressors and inhibitors of the TGF-β signaling pathway; its down-regulation triggers the signal elicited by this factor [63–65]. The fact that spermatozoa from the SPF may carry epididymosomes containing miRNAs in the cauda epididymal fluid [24,66] could explain these findings. These epididymosomes can provide relevant miRNAs to the spermatozoa besides the endogenous ones above listed, potentially enhancing sperm tolerance and female immune tolerance for a successful fertilization and embryo development, influencing fertility. Overall, the current findings contribute to an increased information regarding sperm miRNAs abundance and possible roles of miRNAs via mRNA regulation on the physiological functions and regulatory mechanisms in pig spermatozoa.

Since the breeding boars used in our study were highly selected for sperm quality, an important requisite for incorporation in AI centers, their fertility was far from being a binary variable, and thus it was defined as of higher- or lower- nature, considering farrowing rate (FR) and litter size (LS) as end-points. Consequently, we additionally investigated whether sperm miRNA expression could be somehow related to boar fertility. Increased abundance of mir-182 (SPF) and mir-1285 (post-SRF) and decreased levels of mir-191 (EpiTS) and mir-96 (SRF) were observed in spermatozoa from higher-fertility boars compared with those retrieved from boars with lower fertility. Moreover, since sperm miRNAs could regulate the expression of genes playing important roles in fertilization and embryo developmental ability, as it has been reported before [14], we further investigated the biological implications of the sperm miRNAs we localized and their target genes.

Mir-191 is improperly expressed in some diseases, including cancer, type 2 diabetes, Crohn’s disease, pulmonary hypertension, and Alzheimer’s disease. Nevertheless, little information has been found regarding its role in reproductive processes. mir-191 was proposed as a key factor involved in embryo development. Additionally, higher concentrations of mir-191 in IVF/ICSI medium were observed in non-successful procedures [67]. Interestingly, mir-191 was found within the epididymosomal miRNA content in the bull [40], suggesting to modulate intercellular communication within the epididymis in low-fertility boars, disrupting epididymal function.

The SPF spermatozoa from boars ranked with higher fertility revealed increased levels of mir-182. This finding adds to the findings of Curry et al., 2011, where high expression of mir-182 was observed in boars depicting high sperm motility and intact structure [39]. Many of the identified miRNA target genes are involved in several cellular processes, including cell structure and growth, cellular senescence and calcium signaling, among others. Of particular interest was the influence of mir-182 on the regulation of BCL2L12, IGF1R, MED1, and RARG genes, since these genes are implicated in the promotion of cell
progression and the negative regulation of apoptotic processes, mainly by direct neutralization of caspase-7 (CASP7) and indirect neutralization of caspase-3 (CASP3), which are known to play essential roles in the execution phase of apoptosis, through the phosphoinositide-3 kinase (PI-3K)-Akt and Ras-Raf-MAPK pathways [68,69]. From our specific analysis of differentially expressed genes, we found an overexpression of three target genes for mir-182 (ABHD13, MFAP3, and PCNX) in HF compared to LF in the SPF. Of special interest was the PCNX gene, which has been speculated to play an important role in the testis, related to spermatogenesis [70]. Overall, these results suggest that mir-182 regulates the survival of spermatozoa towards an adequate progression of spermatozoa within the oviduct. mir-96 has been widely involved in the diagnosis of human cancer progression and development because of its involvement in cellular apoptosis and death [71]. Our results indicate that mir-96 was downregulated in the SRF fraction of the boars depicting higher fertility, but the connection with sperm apoptosis in mature pig spermatozoa is still a debated issue, compared to other species (humans for instance, where the degree of chromatin compaction is lower). In consequence, this issue ought to be followed. CACNA2D2, SPEN, and EBF3 are mir-96 target genes found upregulated in spermatozoa from HF compared to LF boars in the SRF. The calcium channel, voltage-dependent, alpha 2/delta subunit 2 (CACNA2D2) regulates calcium current density and is highly expressed in the testis and has been found under expressed in varicocele patients [72]. Mutations of CACNA2D2 have been associated with reduced male fertility in transgenic mice [73]. SPEN (spen family transcription repressor) is a gene that encodes a nucleic acid-binding protein putatively involved in repression of gene expression. SPEN is involved in general downregulation of the transcription during the heat shock response in mouse spermatogenic cells through its interactions with chromatin [74], and methylation changes in EBF3 (Early B cell Factor 3) have been associated with loss of fertility in human [75]. Although the present study revealed some interesting information regarding the abundance of miRNAs in the different sperm sources, it is important to note that other ncsRNAs, such as iRNAs, tsRNAs, or piRNAs are present in spermatozoa, and might be playing an important role in reproductive functions. Further studies are needed to explore potential involvement of other ncsRNAs of interest in sperm function.

5. Conclusions

In conclusion, the results presented in this study revealed novel miRNAs in pig spermatozoa whose relative degrees of abundance vary among spermatozoa, from the functional terminal segment of the epididymis and different fractions of the ejaculate. Some of these specific miRNAs could be linked, for homology to predicted target genes with relevant functions related to sperm survival, immune tolerance, or cell cycle regulation, among others. Such regulation could influence embryo development and, ultimately, fertility of the sires, a matter hereby explored. If these genes are essential, then exploration of sperm miRNAs would be beneficial towards the identification of fertility biomarkers, benefiting the efficiency of artificial insemination techniques through safer selections of the most fertile breeders. However, further research is needed to shed evidence on the mechanistic and physiological roles of such miRNAs, and whether they are intrinsic in spermatozoa or derived from epididymosomes and/or seminal plasma.

Supplementary Materials: The following supporting information can be downloaded at: www.mdpi.com/article/10.3390/biology11020236/s1, File S1. Sample grouping represented in a PCA plot. Information regarding library quality and sequencing outcomes.

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Data Availability Statement: The raw datasets generated during and/or analyzed during the current study are available at Sequence Read Archive (SRA) with BioProject accession number: PRJNA762225 (https://www.ncbi.nlm.nih.gov/sra/PRJNA762225 (accessed on 12 March 2018)).

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