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Testicular expression of long non-coding RNAs is affected by curative GnRHa treatment of cryptorchidism

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Abstract

Background: Cryptorchidism is a frequent endocrinopathy in boys that has been associated with an increased risk of developing testicular cancer and infertility. The condition is curable by combined surgery and hormonal treatment during early pre-pubertal stages using gonadotropin releasing hormone agonist (GnRHa). However, whether the treatment also alters the expression of testicular long non-coding RNAs (lncRNAs) is unknown. To gain insight into the effect of GnRHa on testicular lncRNA levels, we re-analyzed an expression dataset generated from testicular biopsies obtained during orchidopexy for bilateral cryptorchidism.

Results: We identified *EGFR-AS1*, *Linc-ROR*, *LINC00221*, *LINC00261*, *LINC00282*, *LINC00293*, *LINC00303*, *LINC00898*, *LINC00994*, *LINC01121*, *LINC01553*, and *MTOR-AS1* as potentially relevant for the stimulation of cell proliferation mediated by GnRHa based on their direct or indirect association with rapidly dividing cells in normal and pathological tissues. Surgery alone failed to alter the expression of these transcripts.

Conclusion: Given that lncRNAs can cooperate with chromatin-modifying enzymes to promote epigenetic regulation of genes, GnRHa treatment may act as a surrogate for mini-puberty by triggering the differentiation of Ad spermatogonia via lncRNA-mediated epigenetic effects. Our work provides additional molecular evidence that infertility and azoospermia in cryptorchidism, resulting from defective mini-puberty cannot be cured with successful orchidopexy alone.

Keywords: Spermatogenesis, Cryptorchidism, Male infertility, GnRHa, lncRNAs, Antisense lncRNAs, lincRNAs, Mitosis

Résumé

Contexte: La cryptorchidie est une endocrinopathie fréquente chez les garçons. Elle est associée à un risque élevé de cancer des testicules et d'infertilité. La cryptorchidie peut être soignée par une thérapie incluant une intervention chirurgicale et un traitement hormonal par l'agoniste de l'hormone GnRH. Alors que l'effet de la thérapie sur l'expression des ARNm a été analysé, ses conséquences pour la transcription des longs ARNs non codants (ARNlnc) testiculaires restent inconnues. Afin de mieux comprendre les effets du GnRHa sur les concentrations cellulaires des ARNlnc dans le testicule, nous avons analysé des données d'expression d'ARN par séquençage (ARN-Seq) générées en utilisant des biopsies testiculaires obtenues dans le cadre d'une orchidopexie pour cryptorchidie bilatérale.

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Résultats: Nous avons identifié les *ARNlnc EGFR-AS1*, *Linc-ROR*, *LINC00221*, *LINC00261*, *LINC00282*, *LINC00293*, *LINC00303*, *LINC00898*, *LINC00994*, *LINC01121*, *LINC01553*, et *MTOR-AS1* comme potentiellement pertinents pour la stimulation de la prolifération cellulaire induite par le GnRHa. Cette conclusion fait référence à leur association directe ou indirecte avec la croissance et division cellulaire mitotique rapide dans les tissus normaux et pathologiques. Nous constatons également que la chirurgie seule n'a pas d'effet détectable par ARN-Seq sur l'expression de ces ARNlnc.

Conclusion: Étant donné que certains ARNlnc coopèrent avec des enzymes ayant un effet sur la structure chromatinienne et la régulation épigénétique des gènes, le traitement par GnRHa pourrait substituer la mini-puberté en déclenchant la différenciation des spermatogonies Ad par un mécanisme épigénétique qui dépendrait des ARNlnc. Notre travail révèle des nouvelles pistes moléculaires soutenant l'hypothèse que l'infertilité et l'azoospermie associées avec la cryptorchidie sont la conséquence d'une anomalie de la mini-puberté. Cela explique pourquoi une thérapie efficace de cette pathologie ne nécessite pas seulement l'orchidopexie mais aussi un traitement hormonal.

Mots-clés: Spermatogénèse, cryptorchidie, GnRHa, *ARNlnc*, *ARNlnc* antisense, mitose

Introduction

Long non-coding RNAs (lncRNAs) have emerged as key regulators of gene expression in embryonic stem-cell (ESC) self-renewal and differentiation. In ESCs, lncRNAs are regulated at the genetic level by transcription factor binding to lncRNA gene promoters. A major function of lncRNAs is the regulation of specific gene expression at multiple steps, including the recruitment and expression of basal transcription machinery, post-transcriptional modifications, and epigenetics [1]. lncRNAs have also been proposed to play a targeting role by binding to certain methyltransferases and demethylases, and directing them to specific genomic locations. Depending on the biological context, certain methylation events are stably maintained (e.g., methylation involved in inheritance through mitosis of a silenced heterochromatin state), whereas others have to be amenable to change (e.g., when cells differentiate or respond to environmental cues) [2–5]. The so-called natural antisense transcripts (NATs) have been shown to regulate gene expression by affecting transcription and mRNA stability [2–5]. Almost 80% of the mammalian genome is transcribed, and many genomic loci produce RNAs from both sense and antisense DNA strands [6–8], though the functional importance of most of these transcripts is only poorly characterized.

We have previously demonstrated that the presence of type A dark (Ad) spermatogonia in the testis is a marker of low infertility risk (LIR), whereas low or absent levels (below a critical threshold) indicate high infertility risk (HIR) [9, 10]. Treatment with a gonadotropin-releasing hormone agonist (GnRHa, busserelin) enables the Ad spermatogonia population to recover, significantly improving fertility in HIR patients [11]. GnRHa induces a broad transcriptional response, including genes encoding proteins involved in pituitary development, the hypothalamic-pituitary-gonadal axis, and testosterone synthesis [12]. Earlier work focused on protein-coding mRNAs; consequently, nothing is known about the expression of lncRNAs in the treatment of cryptorchidism.

We identified several hundred GnRHa-responsive lncRNAs, which were grouped into long intergenic non-coding RNAs (lincRNAs) and antisense (AS) lncRNAs. We selected candidates on the basis of their expression profiles and then prioritized them for roles in cell growth, differentiation, and disease based on a literature search in PubMed (www.ncbi.nlm.nih.gov/pubmed/). We also included in this search protein-coding genes located upstream or downstream lincRNAs and sense genes overlapping AS-lncRNAs. In addition, we explored the RNA-RNA interaction data available in the RISE database (<http://rise.life.tsinghua.edu.cn>). Finally, we interpreted lncRNA expression signals in the context of protein/RNA profiling data published by the Human Protein Atlas (www.proteinatlas.org) and RNA-sequencing data from HIR/LIR patients [12]. We propose that certain hormone-responsive lncRNAs may play a role in establishing adult spermatogenesis during pre-pubertal stages of development by controlling testicular cell proliferation.

Materials and methods

Study population and biopsy sample collection

The samples used in this study have been described elsewhere [12–14]. A cryptorchid testis is defined as a testis localized outside of the scrotum and incapable of being brought into a stable scrotal position. Sixteen boys with isolated bilateral cryptorchidism who underwent orchidopexy were prospectively included in this study (Fig. 1). The patients had a median age of 18.5 months (range 8–59 months). During the first orchidopexy, biopsies of the ipsilateral testicle were obtained from all patients. Based on histological evaluation, biopsies were categorized into two groups, Ad- (or HIR) and Ad+ (or LIR). The Ad- group included biopsies with no Ad, and the Ad+ group included testes with Ad spermatogonia (Fig. 1). Cryptorchid boys in the Ad-

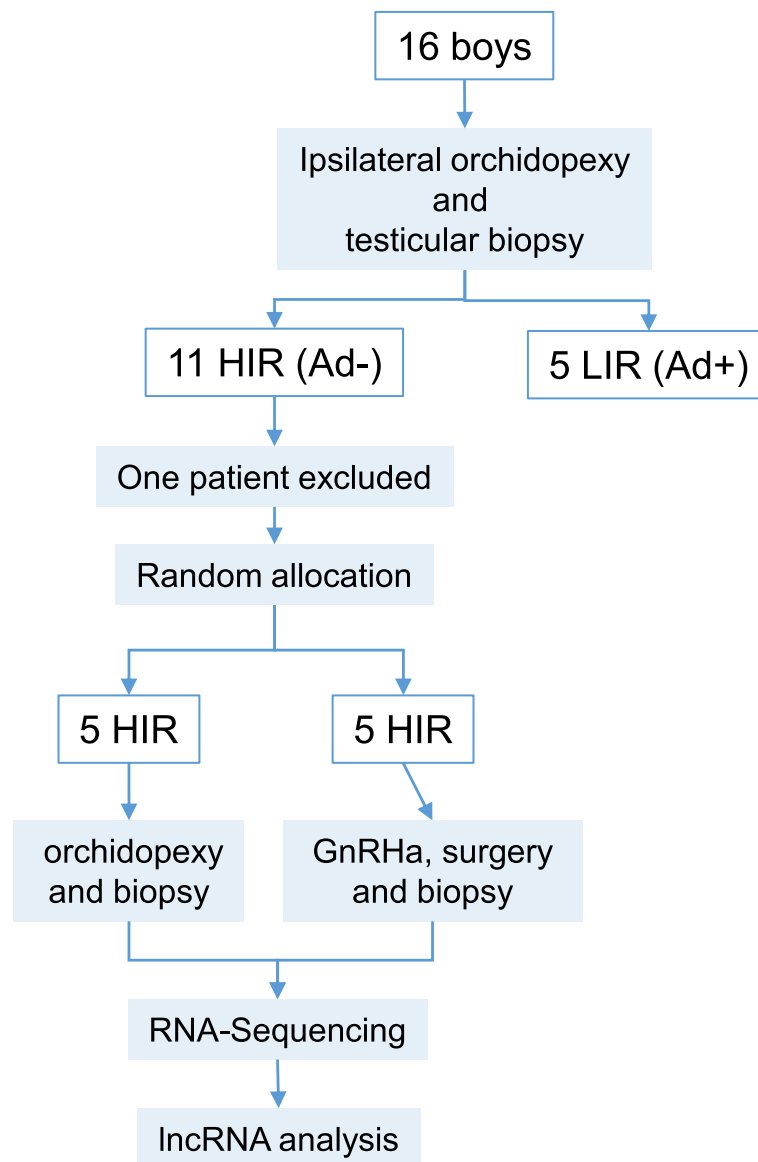


Fig. 1 Flow chart of the study design and the selection of patients and samples for RNA-sequencing based on expression profiling. High infertility risk (HIR) and low infertility risk (LIR) patients are indicated. Classification is based on the absence (Ad-) or presence (Ad+) of Ad spermatogonia

group had 8-times lower plasma LH levels (0.11 IU/L) than the Ad+ group (0.89 IU/L, $p < 0.009$), indicating hypogonadotropic hypogonadism [13]. Five boys (Ad-group) were randomly included in each arm. One HIR patient was excluded (Fig. 1). In the GnRHa-treated group, the median total germ cell count per tubule (S/T) increased from 0.11 to 0.42 ($p = 0.03$, paired-samples Wilcoxon test, one-tailed). In the surgery only group, the median S/T did not change and none of the testes had Ad spermatogonia. In contrast, in the GnRHa treated group, all testes completed the transition from gonocytes to Ad spermatogonia ($p = 0.008$; Fisher test, 2-tailed) [14].

RNA-sequencing data analysis

The workflow from RNA isolation to purification, library preparation, sequencing, data analysis, and expression level analysis has been described in detail elsewhere [12]. Determination of differentially expressed genes, statistical analyses, and model design were carried out as described previously [12]. Genomic coordinates for known lncRNAs were obtained from the Bioconductor package TxDb.Hsapiens.UCSC.hg19.lincRNAsTranscripts (version 3.2.2). Only genes with at least one read per million, in at least two samples were included. P -values and fold-changes were calculated for the treatment factor, and differentially expressed genes were defined as those with a

false discovery rate (FDR) of less than 0.05. Raw data files are available at the Database of Genotypes and Phenotypes (dbGaP) under accession number phs001275.v1.p1. Expression signals are given in standard RKPM units. They are calculated as follows: the number of reads mapped to a gene sequence is divided by the length of the gene sequence over 1000 multiplied by the total number of mapped reads per sample over 1'000'000.

LncRNA data interpretation

We analyzed lincRNA and AS-lincRNA expression in cryptorchid patients with HIR before and after GnRH treatment to identify all RNAs annotated as antisense (AS) transcripts, as well as all RNAs annotated as lincRNAs with $\log_{2}FC > 1.0$. In addition, we compared lincRNA and AS-lincRNA expression between the HIR and LIR groups of cryptorchid patients and analyzed those with lower expression in the HIR group. LincRNAs and AS-lincRNAs were prioritized based on RNA-RNA interactions, revealing the lincRNAs, AS-lincRNAs, or mRNAs encoding proteins involved in spermatogenesis or fertility, and important lincRNAs or mRNAs encoding proteins involved in cell division/growth, signaling pathways, and cancer. Furthermore, we included five lincRNA directly related to spermatogenesis that had a $\log_{2}FC > 1.0$. After the AS-lincRNA candidates were identified, they were prioritized based on a PubMed literature search of themselves and their overlapping sense mRNA/protein, revealing roles in spermatogenesis, fertility, cell division/growth, signaling pathways, and cancer (www.ncbi.nlm.nih.gov/pubmed). The RNA annotation was verified using Ensembl (www.ensembl.org; release 97). The lincRNA/mRNA expression was interpreted using GermOnline (www.germonline.org; version 4.0), Human Protein Atlas (www.proteinatlas.org; version 18), and Genevestigator (www.genevestigator.com; version 7.3.1). Experimentally validated RNA-RNA interaction data were retrieved from RISE (<http://rise.life.tsinghua.edu.cn>; version 1.0).

Results

Global effects on testicular lincRNA levels in response to GnRHa treatment

First, we identified significantly differentially expressed lincRNAs in duplicate testicular biopsies from LIR and HIR patients who underwent surgical correction of undescended testis (Fig. 2, lanes 1–4).

Next, we compared samples obtained from HIR patients at the time of initial surgery (Fig. 2, lanes 5 and 6) and after six months of treatment with GnRHa (Fig. 2, lanes 7 and 8). The genes were ordered using an unsupervised clustering method (hierarchical clustering with complete linkage using Euclidean distances) and are shown in a false-color heatmap relative to the mean expression of each gene over all samples in Fig. 2. The

results indicate that a large number of lincRNAs accumulate at low levels in the testes of boys with HIR compared to LIR, and that a substantial fraction of these transcripts is up-regulated by GnRHa treatment. In contrast, surgery alone had no significant impact on lincRNA expression. We explored the dataset using Volcano plots that display statistical significance (false discovery rate, FDR) against fold-change of expression signals allowing the selection of genes for which large and significant differences in expression levels were observed (Fig. 3).

We found that 627 and 38 lincRNAs were expressed at lower and higher levels, respectively, in HIR versus LIR samples (Fig. 3a). We concluded that the vast majority of differentially expressed lincRNAs are detected at lower levels in HIR testes. Comparing HIR testes before and after GnRHa treatment, we found that 3074 lincRNAs were increased, whereas 53 were decreased (Fig. 3b). Thus, hormonal treatment induces a considerable number of lincRNAs. In the following section the novel lincRNAs were organized based on their known functions or roles that were attributed to their potential target genes.

Certain testicular lincRNAs upregulated by GnRHa treatment are involved in stem cell renewal, signaling, and cell differentiation

We also sought to gain insight into the potential roles that hormone-responsive RNAs might play by interpreting their genomic location, association with protein coding genes in sense/antisense pairs, and RNA-RNA interactions. We selected 11 of 77 lincRNAs and four of 46 AS-lincRNAs with > 2.0 -fold change after GnRHa treatment because their expression patterns lead us to hypothesize that they are important for the development of Ad spermatogonia (Table 1). In this section we focus on novel potential regulatory lincRNAs and we provide context information about their putative protein-coding target genes. This includes previously published expression data obtained with samples from HIR and LIR patients (fold change and FDR values) [12] and functional information relevant for germ cells growth and differentiation from the literature.

LINC01016 is a so-called hub RNA that binds many mRNAs (encoding epigenetic regulators and transcription factors) and lincRNAs (including *XIST*). This feature distinguishes a hub RNA from most other transcripts that interact with few, one or no other RNAs. *LINC01016* is expressed in the same direction as *MLN* (Motilin) and is a transcriptional target of the estrogen receptor [15] (Table 1).

LINC01121 is expressed upstream of *SIX2* and may influence its proximal promoter regions. *SIX2* interacts with *TCF7L2* and *OSR1* in a canonical WNT signaling-independent pathway, preventing the transcription of

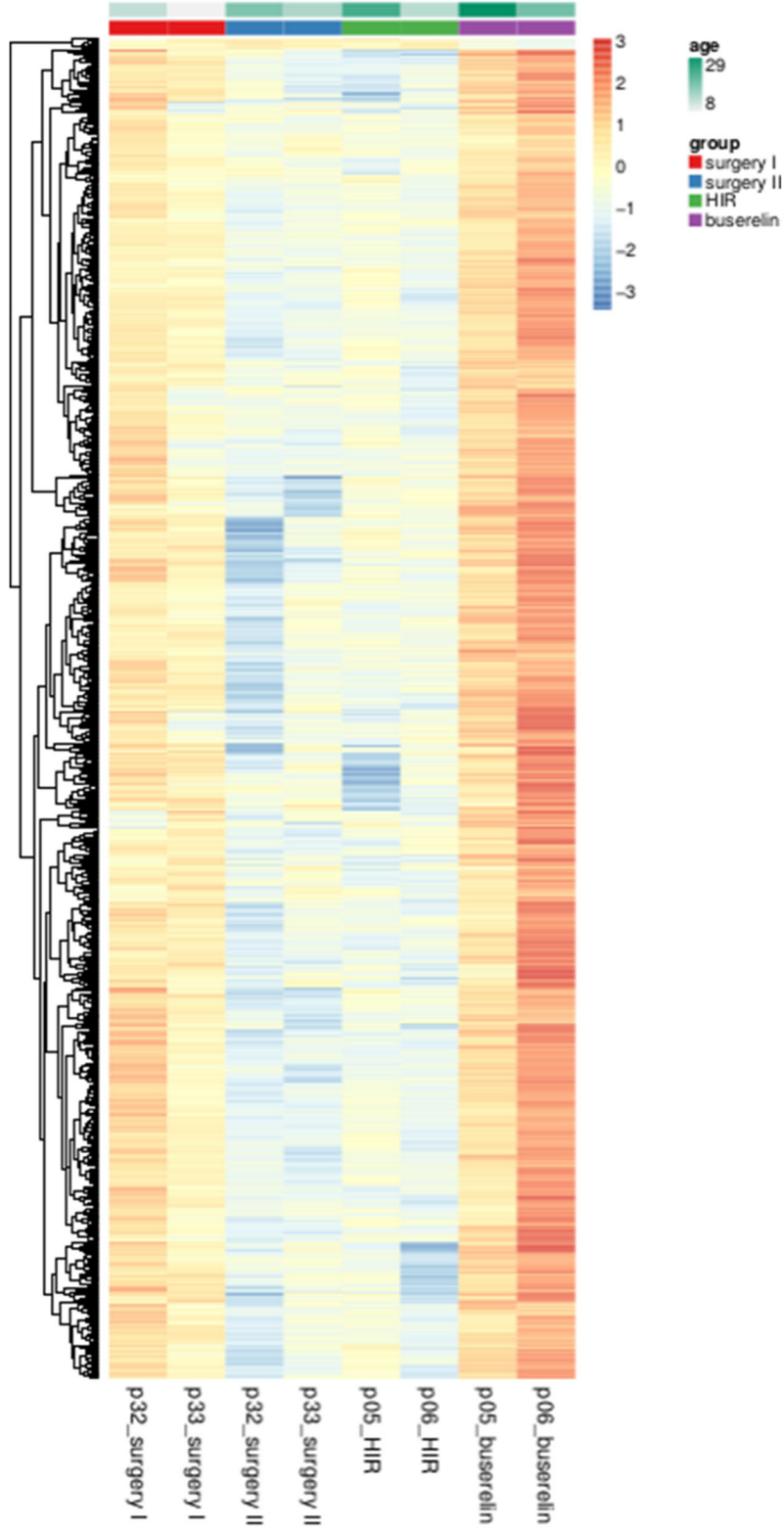


Fig. 2 (See legend on next page.)

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Fig. 2 RNA-sequencing data for lncRNAs. A false color heatmap (red is high, blue is low) shows data from pairs of testes, analyzed. Horizontal bars at the top indicate patient categories and age. Four samples from two HIR patients (p32 and p33; biopsy number) having had surgery “only” treatment; line 1 and 2 during first surgery obtained from ipsilateral testis (I), line 3 and 4 show results from contralateral testis six months after first orchidopexy (II) Buserelin treated patients p5 and p6; line 5 and 6 before treatment and line 7 and 8 contralateral testis after hormonal treatment. Color scales for expression (red/blue) and age (green) are shown

differentiation genes in cap mesenchyme, such as *WNT4* [16–18] (Table 1).

LINC00261 is expressed in the 3' regions of *PAX1* and *FOXA2*, which encode transcription factors. It is also a hub lncRNA that binds mRNAs and lncRNAs, including *HOTAIR*, and is an epigenetically regulated tumor suppressor that is essential for activation of the DNA damage response [19]. *FOXA2* is involved in androgen receptor regulation [19–21] and upregulated after GnRHa treatment ($\log_2FC = 1.69$; $FDR = 0.0004$). Furthermore, *LINC00261* is a tumor suppressor that blocks cellular proliferation by activating the DNA damage response [22].

LINC00303 is expressed upstream of *SOX13*, a developmental factor expressed in mouse Leydig cells and germ cells [23]. Therefore, this lncRNA could be involved in *SOX13* regulation. *LINC00293* is expressed upstream of *SPIDR*, which is involved in double-stranded break repair and genome integrity and binds two TTTY type testis-specific lncRNAs. Several lncRNAs involved in DNA damage repair were increased after GnRHa treatment, including *LINC00994* (expressed

upstream of *PSMD6*), *LINC00898* (binds mRNA encoding *USP1*), and testis-specific *LINC01553* (interacts with mRNA encoding *TIMELESS*). *TIMELESS* plays an important role in the control of DNA replication, the maintenance of genome stability throughout normal DNA replication, and regulation of the circadian clock [24]. (Table 1).

EGFR-AS1, which is involved in determining period length and in the DNA damage-dependent phase advancing the circadian clock [25], interacts with *NEU3* mRNA. *NEU3* activity enhances *EGFR* activation without affecting *EGFR* expression. This may indicate a regulatory mechanism involving a feedback loop. *EGFR-AS1* is weakly expressed in adult testis and highly expressed in liver and liver cancer. Intense *EGFR* immunostaining was found in men with high plasma FSH levels and in all patients who received exogenous FSH, supporting a possible gonadotropin role in the modulation of *EGFR* expression [25]. GnRHa treatment increased the plasma FSH level and *EGFR-AS1*, but decreased *EGFR* expression ($\log_2FC = -0.58$; $FDR = 0.01$). Epidermal growth factor receptor signaling is associated with the pathogenesis of

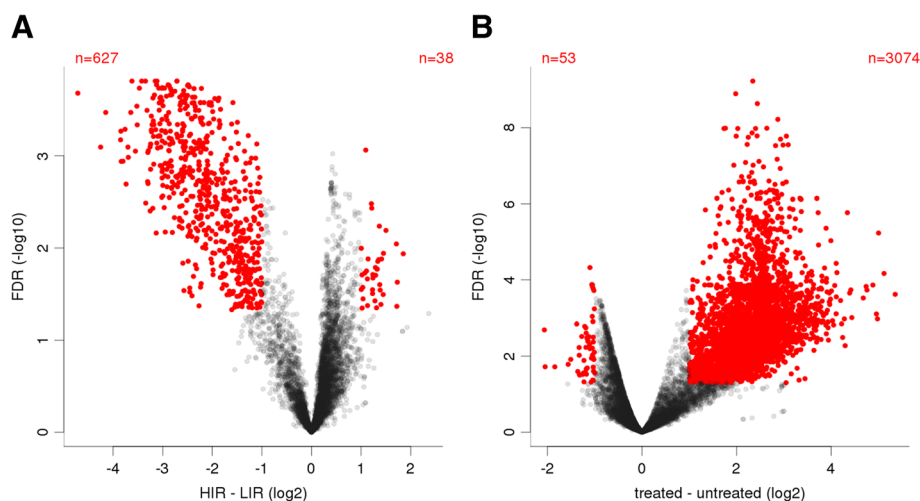


Fig. 3 Volcano plots of lncRNA expression ratios: (a) Between the high (HIR) and low infertility risk (LIR) groups or (b) in the HIR group before and after GnRHa treatment. Genes with no significant difference in expression between the two groups compared in each panel are in black. Differentially expressed genes are shown in red. The most upregulated genes on the right, the most downregulated genes on the left, and the most significant genes at the top

Table 1 Testicular lincRNAs and AS-lincRNAs that increase after GnRHa treatment and are involved in stem cell renewal and differentiation

Gene ID	RPKM before GnRHa Median MAD	RPKM after GnRHa Median MAD	log2FC GnRH	p-value	FDR
<i>LINC-ROR</i>	0.044 / 0.06	0.40 / 0.24	2.68	0.0004	0.002
<i>LINC00261</i>	0.11 / 0.04	1.11 / 0.68	2.60	3.588E-08	4.35E-06
<i>LINC00293</i>	0.05 / 0.04	0.66 / 0.49	2.80	0.0001	0.001
<i>LINC00303</i>	0.22 / 0.07	1.25 / 0.45	2.57	0.0001	0.0008
<i>LINC00520</i>	0.15 / 0.07	0.79 / 0.64	2.76	0.0002	0.001
<i>LINC00898</i>	0.04 / 0.03	0.35 / 0.25	2.72	0.0002	0.001
<i>LINC00974</i>	0.07 / 0.04	0.48 / 0.46	3.9	0.0008	0.003
<i>LINC00994</i>	0.20 / 0.10	1.41 / 0.81	2.73	5.773E-06	0.0001
<i>LINC01016</i>	0.15 / 0.03	1.32 / 0.70	3.36	2.078E-09	6.43E-07
<i>LINC01121</i>	0.25 / 0.04	1.84 / 0.45	2.89	2.049E-09	6.43E-07
<i>LINC01553</i>	0.09 / 0.06	1.26 / 0.67	3.60	0.0002	0.001
<i>EGFR-AS1</i>	0.03 / 0.05	0.58 / 0.30	2.99	3.850E-07	2.28E-05
<i>HOTTIP</i>	0.04 / 0.03	0.47 / 0.37	2.22	0.0010	0.004
<i>MTOR-AS1</i>	0.21 / 0.19	1.90 / 1.33	4.96	6.67E-06	0.0001
<i>OTX2-AS1</i>	0.09 / 0.007	0.91 / 0.45	2.37	2.95E-06	8.83E-05

The log-fold changes (FC), p-value, false discovery rate (FDR), median expression values in reads per kilobase per million (RPKM) (Median), and the median absolute deviation (MAD) for LINC samples before and after GnRHa treatment are given

cutaneous squamous cell carcinoma. *LINC00520*-targeted *EGFR* inhibition might result in inactivation of the PI3K/Akt pathway, thereby inhibiting cancer development [26].

HOTTIP mediates the regulation of *CXCL* genes, which are implicated in Ad spermatogonia differentiation [12]. *HOTTIP* is antisense to *HOXA13* and modulates cancer stem cell properties in human pancreatic cancer by regulating *HOXA9* [27, 28]. *OTX2-AS1* is a NAT RNA that plays an important role in eye development and exhibits sequence complementarity to the exon sequences in its corresponding sense gene, *OTX2*, in both mice and humans [12]. *OTX2* is downregulated in HIR (log2FC = -1.73; FDR = 0.02;) and upregulated after GnRH treatment (log2FC = 1.24; FDR = 0.03) [12]. Though no role has been found for *OTX2-AS1*, deletion of its sense gene *OTX1* was found in six patients with genitourinary defects. Three of these individuals were diagnosed with cryptorchidism [29]. *MTOR*, the key regulator of spermatogenesis [30], is downregulated in boys with HIR (log2FC = -0.42; FDR = 0.03;) and remains downregulated after GnRHa treatment (log2FC = -0.53; FDR = 0.02;). Its antisense gene, *MTOR-AS1*, was up-regulated 4.9-log2 by GnRHa treatment (Table 1). Thus far, nothing is known about the function of *MTOR-AS1*.

LINC-ROR is induced 6.5-fold by GnRHa. (Table 1) This lincRNA controls stem cell renewal and acts as an miRNA sponge via gene silencing, which indicates that the transcript itself has a biological role [31–33]. In addition, we found that *BODIL2*, a testis-specific gene, is located

downstream of *LINC-ROR* and may be transcriptionally regulated by the lincRNA. *BODIL2* plays a role in chromosome biorientation through the detection or correction of syntelic attachments in mitotic spindles [34, 35]. Buserelin treatment increases *BODIL2* gene expression (log2FC = 1.72 l; FDR = 0.003;), indicating a possible role for it in Ad spermatogonia differentiation.

LincRNAs downregulated in HIR testes and stimulated after GnRHa treatment are associated with cancer and the transition of ad spermatogonia

We previously reported different lincRNA expression in patients with HIR compared to LIR; some of these RNAs participate in epigenetic processes, including *AIRN*, *ERICH-AS1*, *FENDRR*, *HAGLR*, and *XIST* [12]. Here, we focus on seven lincRNAs with decreased gene expression in HIR, indicating abrogated mini-puberty, and increased expression after GnRHa treatment (Tables 2 and 3).

LINC00922 is expressed upstream of cadherins *CDH5* and *CDH11*; the latter encodes a calcium-dependent cell adhesion protein that may play a role in testicular architecture [36].

LINC00221 binds mRNA encoding *DCBLD2*, which is involved in negative regulation of cell growth. Significant downregulation of *DCBLD2* occurred after GnRHa treatment (log2FC = -1.0; FDR = 0.0002;). *LINC00221* interacts with *VPS53*'s mRNA. The protein acts as component of the GARP complex involved in retrograde transport from early and late endosomes to the trans-Golgi network [37].

Table 2 Testicular lincRNAs downregulated in HIR testes compared to LIR

lincRNA	LIR Median/MAD	HIR Median/ MAD	log2FC	p-value	FDR
<i>LINC00922</i>	0.47 0.17	0.10 0.07	- 1.47	0.002	0.01
<i>LINC00221</i>	0.64 0.27	0.27 0.07	- 1.20	0.0007	0.008
<i>LINC01249</i>	0.43 0.28	0.15 0.08	- 1.42	0.001	0.01
<i>LINC00701</i>	0.29 0.03	0.11 0.06	- 0.84	0.002	0.02
<i>HOTAIR</i>	0.49 0.23	0.13 0.09	- 1.74	0.0001	0.002
<i>DLX6-AS1</i>	0.42 0.09	0.20 0.06	- 0.92	0.006	0.03
<i>LINC01446</i>	0.57 0.09	0.31 0.08	- 1.21	0.0002	0.003

The log-fold changes (FC), p-value, false discovery rate (FDR), median expression values in reads per kilobase per million (RPKM) (Median), and the median absolute deviation (MAD) for LINC samples are presented

LINC01249 is expressed upstream of *SOX11*, which is important for embryonic neurogenesis and tissue modeling. *SOX11* is upregulated after GnRHa treatment (log2FC = 0.7; FDR = 0.008). It has been suggested that, together with *SOX4*, *SOX11* may function as a transcriptional repressor in fetal testes, contributing to the precise regulation of *SRY* and *SOX9* [23].

LINC01446 promotes glioblastoma progression by modulating the miR-489-3p/*TPT1* pathway [38].

The testis expression of *LINC00701* is developmental stage-specific and associated with *SLC25A37*, encoding a solute carrier localized in the inner mitochondrial membrane. The protein functions as an essential iron importer for the synthesis of mitochondrial heme and iron-sulfur clusters [39].

HOX antisense intergenic RNA (*HOTAIR*) is an lincRNA that coordinates with chromatin-modifying enzymes, regulates gene silencing, and is transcriptionally induced by estradiol (E2) [12, 40, 41].

Distal-less homeobox6 antisense (*DLX6-AS1*) was downregulated in HIR and responded positively to GnRHa treatment. This supports the observation in mice that *DLX6* participates in the control of steroidogenesis [42]. *DLX5* and *DLX6* showed low or no expression in HIR samples.

TINCR, an lincRNA required for the induction of key differentiation genes, is downregulated in HIR testes (log2FC = - 1.07; FDR = 0.002). Seven epigenetic modifiers found to bind *TINCR* were upregulated in HIR and downregulated after GnRHa treatment (Table 4).

Discussion

In this study, we aimed to gain molecular insight into the effect on testicular lincRNA expression levels of a curative treatment for cryptorchidism and related infertility that combines surgery and nasal administration of GnRHa [11, 12, 14]. We found hundreds of lincRNAs that respond to treatment, including a subset that is present at lower levels in testicular samples from boys with HIR. A detailed interpretation of the expression data revealed candidate lincRNAs that may play important regulatory roles in establishing adult spermatogenesis during early postnatal development in humans. Our data are consistent with the hypothesis that hypogonadotropic hypogonadism in boys with altered mini-puberty is the consequence of a profoundly altered gene expression program involving protein-coding genes and lincRNAs. The results point to molecular mechanisms that underlie the ability of GnRHa to rescue fertility.

Study design for human testicular RNA profiling experiments

When working with human samples, a critical issue is the number of cases included in a given analysis. The number of replicates affects the statistical confidence level, and human tissue samples exhibit intrinsic variability that needs to be controlled. In this exploratory lincRNA profiling study, we included first seven patients chosen sequentially from a study based on randomized patient samples [12, 14]. Their inclusion in the cohorts to be treated or to remain untreated was completely unbiased by any parameter other than undescended testes, which were surgically corrected. This sample size, while small, is enough for an initial transcriptome study as presented here.

Table 3 LincRNAs downregulated in HIR testes and stimulated after GnRHa treatment

lincRNA (RPKM)	before GnRHa Median MAD	after GnRHa Median MAD	log2FC	p-value	FDR
<i>LINC00922</i>	0.10 0.07	0.85 0.40	1.41	0.02	0.04
<i>LINC00221</i>	0.27 0.07	1.12 0.42	1.23	0.01	0.03
<i>LINC01249</i>	0.15 0.08	0.92 0.56	1.38	0.003	0.01
<i>LINC00701</i>	0.11 0.06	0.72 0.48	1.28	0.0009	0.003
<i>HOTAIR</i>	0.13 0.09	1.04 0.87	1.14	0.01	0.03
<i>DLX6-AS1</i>	0.20 0.06	1.13 0.81	2.02	1.55E-05	0.0002
<i>LINC01446</i>	0.31 0.08	1.49 0.82	1.14	0.0007	0.003

The log-fold changes (FC), p-value, false discovery rate (FDR), median expression values in reads per kilobase per million (RPKM) (Median), and the median absolute deviation (MAD) for LINC samples before and after GnRHa treatment are shown

Table 4 Seven epigenetic modifiers that bind TINCR

Gene ID	Name	Log2FC HIR/LIR	FDR HIR/LIR	Log2FC HIR/ GnRHa	FDR HIR/ GnRHa
<i>SETD7</i>	SET domain containing lysine methyltransferase 7	+ 0.22	0.042	-0.85	0.0006
<i>ARID4B</i>	AT-rich interaction domain 4B	+ 0.18	0.041	-0.63	0.008
<i>ARID5B</i>	AT-rich interaction domain 5B	+ 0.35	0.004	-0.49	0.003
<i>KDM6A</i>	lysine demethylase 6A	+ 0.20	0.017	-0.81	0.0009
<i>CHD6</i>	chromodomain helicase DNA binding protein 6	+ 0.21	0.040	-0.54	0.03
<i>MBD2</i>	methyl-CpG binding domain protein 2	+ 0.23	0.029	-0.68	0.004
<i>BPTF</i>	bromodomain PHD finger transcription factor	+ 0.19	0.055	-0.61	0.01

The log-fold changes (FC), p-value, false discovery rate (FDR), comparing HIR and LIR cryptorchid testes as well as results from HIR group following GnRHa treatment are presented

Curative hormone treatment affects signaling pathways

During GnRHa treatment, increased LH and testosterone secretion induced the transition of gonocytes and undifferentiated spermatogonia into Ad spermatogonia. In this context, it is interesting that the expression of *LINC-ROR*, a key regulator of pluripotent stem cell reprogramming, increased after hormone treatment. *LINC-ROR* influences cell differentiation, in part, by acting as a sponge for miR-138 and miR-145 and by activating both the canonical and non-canonical WNT/ β -catenin signaling pathways [31]. Importantly, an increase in *mTOR-AS1* expression after GnRHa treatment may have resulted in the suppression of *mTOR* activity, allowing spermatogonial stem cells to undergo self-renewal. *WNT3* induces many transcription factors associated with mesoderm and is downregulated in the testes of men with HIR testes [12]. *WNT* interacts with *mTOR* signaling to affect cancer cell growth and tumor metabolism [43], as well as the formation of spermatozoa [30]. We propose that *WNT3* is the signaling component that regulates early expression of *Brachyury (T)* [44], a mesodermal factor known to determine the fate of Ad spermatogonia. T is a classical and conserved mesodermal factor essential for robust activation of the germline determinants *PRDM1* and *PRDM14* [45]. T directly upregulates these genes, thereby delineating the downstream primordial germ cell program. In mutant mice lacking *Bmp4*, a program induced by *WNT3* prevents T from activating *PRDM1* and *PRDM14*, demonstrating a permissive role of *Bmp4* in primordial germ cell specification [45]. We found that *DMRTC2*, *PAX7*, *T*, and *TERT* are downregulated in testes with defective mini-puberty and respond to GnRHa treatment [46]. Furthermore, we found lower levels of *PRDM1*, *PRDM6*, *PRDM9*, *PRDM13*, and *PRDM14* mRNA in the testes of patients with HIR compared to LIR, and *PRDM7*, *PRDM9*, *PRDM12*, and *PRDM16* were significantly induced after GnRHa treatment. Thus, GnRHa treatment induces an alternate pathway to stimulate *PRDM9* for Ad spermatogonia specification without the permissive role of *BMP4*,

but increased the expression of *BMP5* (log2FC = 2.31; FDR = 0.0001); [12, 47]. *LINC01121* is expressed upstream of *SIX2* and may therefore influence its expression. *SIX2* and *WNT* regulate the self-renewal and commitment of nephron progenitors through shared gene regulatory networks [16–18]. *SIX2* also activates the expression of *GDNF* and plays a role in cell proliferation and migration. Testes in men with HIR expressed lower levels of *SIX2* than LIR testes (log2FC = -1.76; FDR = 0.0008). GnRHa treatment increased *SIX2* (log2FC = 1.61; FDR = 0.014) and *GDNF* (log2FC = 1.46; FDR = 0.003) expression. Taken together, our results support the notion that *LINC-ROR* and *mTOR* are involved in Ad spermatogonia development and specification via the WNT signaling pathway.

Hormone treatment influences epigenetic factors

LINC00261 stimulates the expression of *HOTAIR* and *HOTTIP* genes to stabilize androgen receptor (AR) together with *FOXA1*. Specifically, the DNA-binding domain (DBD)/hinge region of AR directly interacts with the fork head domain of *FOXA1*, thereby acting as an AR-collaborating factor [48].

HOTAIR's promoter contains multiple functional estrogen response elements (EREs). *HOTAIR* mediates the recruitment of H3K27 methyltransferase and H3K4 demethylase, which leads to efficient repression of certain loci. The levels of histone H3 lysine-4 trimethylation, histone acetylation, and RNA polymerase II recruitment are increased at the *HOTAIR* promoter in the presence of E2, and knock-down of ERs downregulated E2-induced *HOTAIR* expression. Thus, like the transcription of protein-coding genes, E2 induces the transcription of antisense RNAs [27, 49, 50]. HIR patient's exhibit decreased *HOTAIR* levels, [12] and GnRHa treatment induced *HOTAIR* expression, indicating that LH, testosterone and/or converted E2 have a positive effect on *HOTAIR* and, thus, Ad spermatogonia differentiation [12]. Furthermore, *HOTAIR* was previously shown to mediate tumorigenesis by recruiting *EZH2* [51]. This is of interest, as GnRHa induces expression of *HOTAIR*, downregulates *EZH2* (log2FC = -0.6; FDR = 0.01) and

implicates the regulation of *HOTTIP* gene transcription, which then transcriptionally regulates the *HOXA* cluster. As a result, an increase occurs in *HOXA2* (log2FC = 2.38; FDR = 0.007), *HOXA3* (log2FC = 1.68; FDR = 0.007), *HOXA11* (log2FC = 1.77; FDR = 0.02), and *HOXA-AS3* (log2FC = 2.68; FDR = 0.0001).

Gamma-aminobutyric acid (GABA) plays a key developmental role in the regulation of GnRH neuron migration from the olfactory placodes into the forebrain during fetal development [52], and co-expression of *DLX3* and *PAX6* proteins, correlates with acquisition of the olfactory placode fate [53]. Moreover, *GABA-A* receptors and GABA transporter 1 (*GAT1*) have been reported to be involved in the proliferation of Leydig cells, testosterone production, and spermatogenesis [54]. *GABRA 3* (log2FC = -2.49; FDR = 0.0001;) and *GABR5* (log2FC = -2.59; FDR = 0.0006;) were downregulated in HIR testes. Following GnRHa treatment, an increase expression was observed in *DLX3*, *PAX6* [12], and *TP63* (log2FC = 0.91; FDR = 0.002;), whereby the latter was downregulated in HIR (log2FC = -1.58; FDR = 0.001;). Berghoff et al. proposed a model in which *DLX6-AS1* inhibits the ultraconserved DNA methylation mark in *DLX5/6*, facilitating antagonistic interactions between repressive and activating transcription factors *MECP2* and *DLX* [53]. *DLX2*, *DLX3*, *DLX5* and *DLX6* showed low or no expression in HIR samples [12] Loss of *DLX1/2* increases site-specific methylation of the *DLX5/6* ultraconserved enhancer [53]. *TP63* regulates *DLX5* and *DLX6* transcription, at least in part, via cis-acting regulation at the promoter level [55]. These interactions allow differential control of adjacent genes by shared DNA regulatory elements [56]. It was also shown that deletion of *Dlx5* and *Dlx6* in the mouse leads to decreased testosterone levels and an abnormal masculinization phenotype [42]. Thus, impaired steroidogenesis during mini-puberty in HIR boys may be induced by altered *GABRA-A* receptor signaling, silenced *DLX6* expression as well as destabilization of AR and ERs.

GnRHa treatment affects genes associated with normal and pathological cell division

Several recent studies have reported roles for lncRNAs in cancer, including *MALAT1* and *GAS*, strongly indicating that lncRNAs not only control gene regulatory pathways in normal cells and tissue, but also during tumor development [57, 58]. The occurrence and progression of cancer is the result of a combination of multiple factors. Furthermore, cancer-specific lncRNA expression patterns appear to be more tissue- and stage-specific than those of protein-coding genes, supporting the potential development of lncRNAs as efficient biomarkers and therapeutic targets [58]. Interestingly, GnRHa treatment increased LH and testosterone secretion and

HOTAIR expression, and probably its recruitment to chromatin, whereas the expression of *MALAT1* was reduced (log2FC = -0.64; FDR = 0.03; REF), suggesting opposite regulation and functions of these lncRNAs during normal testis development. It was reported that LINC-ROR promote liver cancer stem cell growth by upregulating *TERT* and *C-MYC* [59]. Notably, GnRHa treatment stimulates the expression of *TERT* (log2FC = 0.91; FDR = 0.002;) [46], which is decreased in HIR (log2FC = -1.58; FDR = 0.001;) [46] but downregulates *C-MYC* signaling (log2FC = -0.58; FDR = 0.01;).

Variation in lincRNA expression may be associated with cancer progression. For example, *LINC00261*, which plays an important role in gastric cancer, is stimulated by GnRHa, and exerts tumor suppressive activity by reducing cancer cell invasion via suppression of the epithelial-mesenchymal transition process [60]. Another lincRNA, *TINCR*, is involved in normal tissue differentiation and plays a critical role in cancer and metastasis. *TINCR* expression is downregulated in HIR (log2FC = -1.07; FDR = 0.002;) [12]. We observed that *TINCR* depletion in HIR testes resulted in the induction of key epigenetic modifiers, seven of which were downregulated following GnRHa treatment (Table 4). One of them, *SETD7*, is an epigenetic modifier and regulates AR [61]. *SETD7* binds *TINCR* and may mediate the formation of AR-associated coactivator complexes. Taken together, the results indicate that the HIR group of cryptorchid boys with abortive mini-puberty expresses several cancer genes. Gonadotropin-releasing hormone treatment may protect against testicular tumor development by downregulating oncogenes, such as *MALAT1*, *mTOR*, *C-MYC*, and *EZH2* to enable normal germ cell development.

Conclusions

Two major goals in the field of male reproductive biology are to elucidate the molecular mechanisms that underlie cryptorchidism and to develop an effective treatment for its long-term consequences. According to the mini-puberty hypothesis, early-life exposure to gonadal hormones during a specific window of sensitivity triggers sex-specific developmental processes. To preserve molecular features of differentiated cells, it is crucial that transcriptional alterations triggered by external or intrinsic signals continue beyond the initial stimulus. One possible mechanism involves epigenetic histone variant replacement, chromatin remodeling factors, and lncRNAs associated with epigenetic factors. We found that many of the lincRNAs responding to GnRHa treatment are associated with somatic cancer. This may reflect the fact that the hormone stimulates germ stem cell growth and Leydig, as well as Sertoli, cell division. The mechanisms involved are likely rather diverse and may include promoter/enhancer activity, miRNA sponge

activity, or control of gene expression via RNA-RNA interactions. We propose that the lncRNAs identified in this study may be involved in establishing normal male fertility by acting at early stages of spermatogonia stem cell development and by affecting other testicular cells capable of responding to GnRH treatment. Our results provide information's for further functional analysis of long-noncoding RNA in relation to the infertility development. We propose that the HOTAIR and DLX pathways, as well as both canonical and non-canonical WNT pathways, are involved in Ad spermatogonia growth and differentiation.

Abbreviations

1PRDM6: PR/SET domain 6; AIRN: Antisense of IGF2R (insulin like growth factor 2 receptor) non-protein coding RNA; AR: Androgen receptor; ARID4B: AT-rich interaction domain 4B; ARID5B: AT-rich interaction domain 5B; AS-lncRNA: Antisense long non-coding RNA; Bmp4: Bone morphogenetic protein 4; BMP5: Bone morphogenetic protein 5; BOD1L2: Biorientation of chromosomes in cell division 1 like 2; BPTF: Bromodomain PHD finger transcription factor; CDH11: Cadherin 11; CDH5: Cadherin 5; CDH6: Cadherin 6; C-MYC: v-myc avian myelocytomatosis viral oncogene homolog; CXCL: Chemokine; DBD: DNA-binding domain; dbGaP: Database of Genotypes and Phenotypes; DCBLD2: Discoidin, CUB and LCCL domain containing 2; DLX: Distal-less homeobox; DLX2: Distal-less homeobox 2 [*Homo sapiens* (human)]; DLX3: Distal-less homeobox 3 [*Homo sapiens* (human)]; *Dlx5*: Distal-less homeobox 5 [*Mus musculus* (house mouse)]; *Dlx6*: Distal-less homeobox 6 [*Mus musculus* (house mouse)]; DLX6-AS1: Distal-less homeobox 6 antisense RNA 1; DMRTC2: DMRT (doublesex and mab-3 related transcription factor) like family C2; E2: Estradiol; EGFR: Epidermal growth factor receptor; EGFR-AS1: Epidermal growth factor receptor antisense RNA 1; EREs: Estrogen response elements; ERICH-AS1: Glutamate rich protein antisense RNA 1; ERs: Estrogen receptors; ESC: Embryonic stem-cell; EZH2: Enhancer of zeste 2 polycomb repressive complex 2 subunit; FDR: False discovery rate; FENDRR: FOXF1 (forkhead box F1) adjacent non-coding developmental regulatory RNA; FOXA1: Forkhead box A1; FOXA2: Forkhead box A2; FSH: Follicle stimulating hormone; GABA: Gamma-aminobutyric acid; GABA-A: Gamma-aminobutyric acid type A; GABRA-A: gamma-aminobutyric acid (GABA) A receptor, alpha 1; GABRA-A5: Gamma-aminobutyric acid (GABRA) A receptor 5; GARP: Golgi-associated retrograde protein; GAS: Growth arrest specific; GAT1: GABA (gamma-aminobutyric acid) transporter 1; GDNF: Glial cell derived neurotrophic factor; GnRHa: Gonadotropin releasing hormone antagonist; H3K27: Histone H3 Lysine 27; H3K4: Histone H3 Lysine 4; HAGLR: HOXD (homeobox D cluster) antisense growth-associated long non-coding RNA; HIR: High infertility risk; HOTAIR: HOX (homeobox) transcript antisense RNA; HOTTIP: HOXA (homeobox A cluster) distal transcript antisense RNA; HOXA11: Homeobox A11; HOXA13: Homeobox A13; HOXA2: Homeobox A2; HOXA3: Homeobox A3; HOXA9: Homeobox A9; HOXA-AS3: HOXA (Homeobox A) cluster antisense RNA 3; KDM6A: Lysine demethylase 6A; LH: Luteinizing hormone; LINC00221: Long intergenic non-protein coding RNA 221; LINC00261: Long intergenic non-protein coding RNA 261; LINC00282: long intergenic non-protein coding RNA 282; LINC00293: Long intergenic non-protein coding RNA 293; LINC00303: Long intergenic non-protein coding RNA 303; LINC00520: Long intergenic non-protein coding RNA 520; LINC00701: Long intergenic non-protein coding RNA 701; LINC00898: Long intergenic non-protein coding RNA 898; LINC00922: Long intergenic non-protein coding RNA 922; LINC00974: Long intergenic non-protein coding RNA 974; LINC00994: Long intergenic non-protein coding RNA 994; LINC01016: Long intergenic non-protein coding RNA 1016; LINC01121: Long intergenic non-protein coding RNA 1121; LINC01249: Long intergenic non-protein coding RNA 1249; LINC01446: Long intergenic non-protein coding RNA 1446; LINC01553: Long intergenic non-protein coding RNA 1553; lincRNA: Long intergenic non-protein coding RNA; Linc-ROR: Long intergenic non-protein coding RNA, regulator of reprogramming; LIR: Low infertility risk; lncRNA: Long non-coding RNA; log2FC: log2 fold change; MALAT1: Metastasis associated lung adenocarcinoma transcript 1; MBD2: Methyl-CpG binding domain protein 2; MECP2: methyl-CpG binding

protein 2; miR-138: microRNA 138; miR-145: microRNA 1; miR-489-3P: microRNA 489; MLN: Motilin; MTOR: mechanistic target of rapamycin kinase; MTOR-AS1: MTOR (mechanistic target of rapamycin kinase) antisense RNA 1; NAT RNA: N-acetyltransferase RNA; NATs: Natural antisense transcripts; NEU3: Neuraminidase 3; OSR1: Odd-skipped related transcription factor 1; OTX1: Orthodenticle homeobox 1; OTX2: Orthodenticle homeobox 2; OTX2-AS1: Orthodenticle homeobox 2 antisense RNA 1 (head to head); PAX1: Paired box 1; PAX6: Paired box 6; PAX7: Paired box 7; PI3K/Akt pathway: Phosphatidylinositol 3-kinase, putative, and protein kinase B pathway; PRDM1: PR domain containing 1, with ZNF domain; PRDM12: PR/SET domain 12; PRDM14: PR/SET domain 14; PRDM16: PR/SET domain 16; PRDM7: PR/SET domain 7; PRDM9: PR/SET domain 9; PRMD13: PR/SET domain 13; PSMD6: Proteasome 26S subunit, non-ATPase (adenosine triphosphatase) 6; RPKM: Reads per kilo base per million mapped reads; S/T: Total germ cell count per tubule; SETD7: SET domain containing 7, histone lysine methyltransferase; SIX2: SIX homeobox 2; SLC25A37: Solute carrier family 25 member 37; SOX11: SRY (sex determining region Y)-box transcription factor 11; SOX13: SRY (sex determining region Y)-box transcription factor 13; SOX4: SRY (sex determining region Y)-box transcription factor 4; SPDR: Scaffold protein involved in DNA repair; SRY: Sex determining region of Y; T: Brachyury; TCF7L2: Transcription factor 7 like 2; TERT: Telomerase reverse transcriptase; TIMELESS: Timeless circadian regulator; TINCR: TINCR (Tissue differentiation-inducing non-protein coding RNA) ubiquitin domain containing; TP63: Tumor protein p63; TPT1: Tumor protein, translationally-controlled 1; TTTY: Testis-specific transcript, Y-linked; USP1: Ubiquitin specific peptidase 1; VPS53: VPS53 (Vacuolar protein sorting 53 homolog) subunit of GARP (Golgi-associated retrograde protein) complex; Wnt: Wingless-related integration site; WNT3: Wnt family member 3; WNT4: Wnt family member 4; XIST: Inactive X specific transcripts

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Authors' contributions

FH conceived and designed the study, interpreted the data, and organized and wrote the manuscript. GV performed experiments, analyzed the data, and read the paper. BV performed experiments and read the paper. MBS analyzed and interpreted the data, contributed analysis tools, and read the paper.

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Availability of data and materials

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Ethics approval and consent to participate

Investigations were carried out in accordance with the Declaration of Helsinki of 1975 (revised in 2008). The study was approved by the Institutional Review Board and the Independent Ethics Committee of Vilnius University (Vilnius Regional Biomedical Research Ethics Committee, No. 158200–580-PPI-17, 11 June 2013).

Consent for publication

Written informed consent was obtained from the patients' guardians after approval by the ethical committee.

Competing interests

The authors declare no conflicts of interest.

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