Regulation of adrenal and ovarian steroidogenesis by miR-132

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Abstract

miR-132 is hormonally regulated in steroidogenic cells of the adrenal gland, ovary and testis. Here, we examined the potential role of miR-132 in the control of steroidogenesis. Transfection of Y1 adrenal cells with miR-132 increased mRNAs of 3 β -HSD and 20 α -HSD enzymes, which catalyze the sequential conversion of pregnenolone to progesterone to biologically inactive 20 α -hydroxyprogesterone (20 α -OHP). Overexpression of miR-132 reduced MeCP2 and StAR protein expression, basal progesterin (progesterone and 20 α -OHP) production, but enhanced their production in response to cAMP stimulation. Use of [³H] pregnenolone and free-diffusible 22(R)-hydroxycholesterol further confirmed that miR-132 promotes the production of 20 α -OHP by upregulating 3 β -HSD and 20 α -HSD. Evidence is also presented that StAR is a direct target of miR-132. Transient transfection of Y1 cells with miR-132 demonstrated that miR-132 induction of 3 β -HSD and 20 α -HSD was accompanied by significant suppression of one of its target gene products, MeCP2. In contrast, co-expression of miR-132 plus MeCP2 protein partially blocked the ability of miR-132 to upregulate the expression and function of 3 β -HSD and 20 α -HSD. Moreover, suppression of MeCP2 protein with siRNA resulted in increased expression of 3 β -HSD and 20 α -HSD, further demonstrating that miR-132 induces the expression of these two enzymes via inhibition of MeCP2. Likewise, overexpression of miR-132 increased 20 α -OHP production with and without HDL loading, while knockdown of miR-132 resulted in a significant decrease of 20 α -OHP production by granulosa cells. In conclusion, our data suggest that miR-132 attenuates steroidogenesis by repressing StAR expression and inducing 20 α -HSD via inhibition of MeCP2 to generate a biologically inactive 20 α -OHP.

Key Words

- ▶ steroidogenesis
- ▶ StAR
- ▶ microRNA
- ▶ miR-132
- ▶ 20 α -hydroxyprogesterone

*Journal of Molecular
Endocrinology*
(2017) **59**, 269–283

Introduction

Steroid hormones mediate a wide variety of vital physiological functions, including regulation of carbohydrate metabolism, inflammation and stress responses (glucocorticoids), maintenance of blood pressure and water and salt balance (mineralocorticoids) and events connected with fertility and reproduction

(gonadal steroids) (Graham & Clarke 1997, Hiort *et al.* 1998, Newton 2000, Katzenellenbogen *et al.* 2002, Pearce *et al.* 2003). They are mainly synthesized in the adrenal cortex, ovary, testis and placenta. The synthesis of these steroid hormones (steroidogenesis) involves multiple pathways, as well as sequential actions of steroidogenic

enzymes associated with the endoplasmic reticulum (ER) and mitochondria (Jefcoate *et al.* 1992, Stocco & Clark 1996, Stocco 2000, Kraemer & Shen 2002, Payne & Hales 2004, Miller 2008, LaVoie & King 2009, Hu *et al.* 2010, Miller & Bose 2011, Papadopoulos & Miller 2012). Adrenal, ovarian and testicular steroidogenesis is primarily controlled by cell/tissue-specific trophic hormones (ACTH, LH and FSH) (Stocco & Clark 1996, LaVoie & King 2009, Hu *et al.* 2010, Miller & Bose 2011) and is subject to both acute (Jefcoate *et al.* 1992, Stocco & Clark 1996, Stocco 2000, Miller 2008, LaVoie & King 2009, Hu *et al.* 2010, Miller & Bose 2011, Papadopoulos & Miller 2012) and chronic regulation (Simpson *et al.* 1992, Payne & Youngblood 1995, Payne & Hales 2004, Sewer *et al.* 2007, Miller 2008, LaVoie & King 2009, Miller & Bose 2011).

Acute steroid synthesis that occurs over minutes in response to trophic hormone stimulation is controlled at the level of delivery of substrate cholesterol from the outer mitochondrial membrane (OMM) to the inner mitochondrial membrane (IMM), where cytochrome P450 cholesterol side-chain cleavage (P450_{scc}) enzyme (encoded by the *CYP11A1* gene) resides (Jefcoate *et al.* 1992, Stocco & Clark 1996, Stocco 2000, Hu *et al.* 2010, Miller & Bose 2011). This step is primarily mediated by the steroidogenic acute regulatory protein (StAR). Trophic hormones rapidly induce the expression of StAR in their respective target steroidogenic cells (with the exception of steroidogenic cells of the placenta), which then facilitates the transfer of cholesterol from OMM to IMM and thereby initiates steroidogenesis (Stocco & Clark 1996, Stocco 2000, Christenson & Strauss III 2000, Miller & Bose 2011). Chronic stimulation (hours to days) occurs through the trophic hormone induction of P450 enzymes such as *CYP11A*, *CYP11B*, *CYP17*, *CYP19* and *CYP21* and β HSDs, resulting in increased steroidogenic capacity of the cells (Simpson *et al.* 1992, Payne & Youngblood 1995, Payne & Hales 2004, Sewer *et al.* 2007, Miller 2008, LaVoie & King 2009, Miller & Bose 2011).

Although current evidence suggests that both acute and chronic phases of steroidogenesis are mainly regulated at the gene transcription level (Simpson *et al.* 1992, Payne & Youngblood 1995, Payne & Hales 2004, Sewer *et al.* 2007, Miller 2008, LaVoie & King 2009, Manna *et al.* 2009, Miller & Bose 2011), emerging evidence also suggests that steroidogenesis may be subject to posttranscriptional and posttranslational regulations (Strålfors *et al.* 1984, Epstein & Orme-Johnson 1991, Arakane *et al.* 1997, Greenberg *et al.* 2001, Nair *et al.* 2002, Lehoux *et al.* 2004, Manna *et al.* 2009, Wang *et al.* 2009, Hu *et al.* 2012, 2013a,

2015, Kitahara *et al.* 2013, Menon *et al.* 2013, Morohaku *et al.* 2014). An earlier publication from our laboratory showed that scavenger receptor class B, type 1 (SR-B1), an HDL receptor that mediates bulk delivery of HDL-derived CE into the steroidogenic cells of the adrenal gland and ovary and to testicular Leydig cells under certain conditions (Azhar *et al.* 1998, 2002, Reaven *et al.* 2002, Hu *et al.* 2010), is also subject to posttranslational regulation by Na⁺/H⁺ exchanger regulatory factors 1/2 (NHERF1/2) (Hu *et al.* 2013a). In addition, we recently provided evidence that miRNAs, such as miR-125a and miR-455, target SR-B1 mRNA, thus posttranscriptionally and negatively regulate SR-B1-mediated selective delivery of HDL cholesterol in steroidogenic cells and consequently inhibit SR-B1-mediated and HDL-supported steroidogenesis (Hu *et al.* 2012). In contrast, we very recently provided evidence that SIK1 protein kinase positively regulates HDL-CE transport function of SR-B1 and secondarily steroidogenesis (Hu *et al.* 2015).

MicroRNAs (miRs) comprise a novel class of non-coding RNA single-stranded small RNA transcripts of 20–25 nucleotides that regulate gene expression at the posttranscriptional level. They have emerged as key posttranscriptional regulators of gene expression in metazoan animals, plants, protozoa and viruses (Ambros 2004, Jones-Rhoades *et al.* 2006, Bushati & Cohen 2007, Bartel 2009, Fabian *et al.* 2010, Skalsky & Cullen 2010). Recently, we reported the high-throughput expression profiling of miRNAs in rat adrenals in response to *in vivo* treatment of animals with hormones, ACTH, 17 α -ethinyl estradiol (17 α -E2) or dexamethasone (Hu *et al.* 2013b). Among the various adrenal miRNAs whose expression was altered in response to treatment of rats with any of these three hormones, the most robust effect was on the expression of miR-132 and miR-212 (3–4 folds' induction) in response to ACTH treatment (Hu *et al.* 2013b). We further demonstrated that the expression of both miR-132 and miR-212 was upregulated in primary rat ovarian granulosa cells and mouse testicular Leydig tumor cells by the second messenger of the tropic hormone, cAMP (Hu *et al.* 2013b). These findings, coupled with other reports (Fiedler *et al.* 2008, Yao *et al.* 2010), suggest the possibility that these two miRNAs may participate in the trophic hormone regulation of steroidogenesis. miR-132, together with miR-212, comprises the evolutionary conserved miR-132/212 family, encoded from the same intron of a small non-coding gene that is located on chromosome 11 in mice, chromosome 10 in rats and chromosome 17 in humans. Mature miRNA-132/-212 share the same seed sequence (Wanet *et al.* 2012, Remenyi *et al.* 2013).

Here, we examined the potential role of miR-132 in the regulation of steroidogenesis in a model adrenal cell line, Y1 cells. Given that miR-212 expression was several hundredfold lower, as compared to miR-132, in all of the steroidogenic cells examined here, we focused our studies primarily on miR-132. Our findings suggest that miR-132 inhibits steroidogenesis by causing the direct repression of StAR protein expression and by promoting the increased conversion of biologically active progesterone into biologically inactive 20 α -hydroxyprogesterone (20 α -OHP). The latter effect of miR-132 is mediated by increased expression of 20 α -hydroxysteroid dehydrogenase (20 α -HSD) via the miR-132 inhibition of methyl-DNA-binding protein, MeCP2; 20 α -HSD catalyzes the conversion of progesterone into its inactive form, 20 α -OHP (Mao *et al.* 1997).

Materials and methods

Materials

22(R)-hydroxycholesterol, 17 β -estradiol, N6,2'-O-dibutyryl adenosine 3',5'-cyclic monophosphate (Bt₂cAMP), progesterone, 20 α -OHP, 22(R)-hydroxycholesterol and fatty acid-free bovine serum albumin were supplied by Sigma-Aldrich. 20 α -hydroxyprogesterone [1,2-³H(N)] (40–60 Ci/mmol; 1.48–2.22 TBq/mmol) and progesterone [1,2-³H] (40–60 Ci/mmol; 1.48–2.22 TBq/mmol) were purchased from American Radiolabeled Chemicals. Pregnenolone, [7-³H(N)] (10–25 Ci/mmol; 370–925 GBq/mmol) was supplied by PerkinElmer. StAR (D10H12) XP rabbit mAb, GAPDH (14C10) rabbit mAb and MeCP2 (D4F3) XP rabbit mAb were obtained from Cell Signaling Technology. Purified anti-SF-1 polyclonal antibody (07-618) was purchased from EMD Millipore. All other chemicals used were of analytical grade.

Animals and hormonal treatment

All animal experiments were performed according to the procedures approved by the VA Palo Alto Health Care System Animal Care and Use Committee (IACUC). Groups of eight 225- to 250-g male Sprague–Dawley rats were purchased from Harlan Laboratories (Indianapolis, IN, USA). They were allowed to acclimatize to a new controlled environment (25 $\pm$ 2°C, 55% $\pm$ 5% relative humidity with a 12-h light-darkness cycle) for approximately 1 week. Subsequently, the rats were randomly assigned to one of two groups (4 rats in each group): group 1 rats were treated subcutaneously with phosphate-buffered saline

(PBS) every 24 h for 4 days, with the last injection on day 4 given 1 h prior to harvesting of adrenal tissues, and the rats in group two were treated subcutaneously with ACTH (Cortrosyn) (10 IU) every 24 h for 4 days, and again, the ACTH injections on day 4 given 1 h prior to killing the animals. The collected adrenals were stored frozen in liquid nitrogen until their use in RNA isolation. Similarly, immature female Sprague–Dawley rats (21–23 days old, Harlan Laboratories) were injected subcutaneously with 17 β -estradiol (1 mg/kg BW) daily for 5 days. The animals were killed 24 h after their last injection, and the granulosa cells were isolated from the ovaries using the follicle puncture method as described previously (Azhar *et al.* 1998).

Isolation, culture and cAMP treatment of rat ovarian granulosa cells

Isolated granulosa cells (Azhar *et al.* 1998) were cultured in 35-mm culture dishes that were precoated with 1% fetal bovine serum overnight. Dishes were plated with 1–2 $\times$ 10⁵ cells in Dulbecco's modified Eagle's (DME)-F12 culture medium containing 15 mM Hepes, 1 mg/mL bovine serum albumin, 2 μ g/mL insulin, 5 μ g/mL transferrin, 100 ng/mL hydrocortisone, 100 μ g/mL streptomycin and 100 U/mL penicillin G. After 72 h of culture, cells were treated with or without Bt₂cAMP (2.5 mM) for 24 h and subsequently used for experimentation.

Cell culture

Y1 mouse adrenocortical cells (ATCC CCL-79), MLTC-1 mouse Leydig tumor cells (ATCC CRL-2065) and CHO-K1 cells (ATCC CCL-61) were purchased from the American Type Culture Collection (ATCC). Y1 cells were cultured in Ham's F-12K (Kaigh's) medium containing 2.5% fetal bovine serum, 15% horse serum and 100 units/mL penicillin and 100 μ g/mL streptomycin. In contrast to mouse adrenocortical cells/adrenal gland that produce corticosterone as the major steroid product, cultured Y1 cells, both under basal conditions and in response to trophic hormone stimulation, synthesize and secrete 20 α -hydroxy- Δ^4 -pregnene-3-one (20 α -hydroxyprogesterone) and 11 β ,20 α -dihydroxy- Δ^4 -pregnene-3-one (11 β , 20 α -hydroxyprogesterone) (Rainey *et al.* 2004). Y1 cells also secrete small quantities of progesterone. The lack of production of a normal steroid, corticosterone, by these cells is mainly attributed to CYP21 (steroid 21-hydroxylase) deficiency (Parker *et al.* 1985). MLTC-1 cells were cultured in

RPMI-1640 medium supplemented with 10% fetal bovine serum plus 100 U/mL penicillin and 100 µg/mL streptomycin. These cells mainly secrete progesterone and a small amount of testosterone (Manna *et al.* 2006). CHO-K1 cells were cultured in F-12K medium containing 10% fetal bovine serum and 100 units/ml penicillin and 100 µg/mL streptomycin. All cell cultures were maintained at 37°C in a tissue culture incubator containing 5% CO₂, 95% air. Most of the tissue culture supplies were purchased from Invitrogen through Invitrogen Cell Culture Media.

Plasmid constructs

The open reading frame of mouse steroidogenic acute regulatory protein (mStAR) was amplified and subcloned into pcDNA6-HisA-V5 vector, and the constructed

plasmid was designated as pcDNA6-mStAR-V5. The mouse *StAR* (166bp (562–727 nucleotides (nt) from the start of the 3'UTR) and 213bp (1185–1397 nt from the start of the 3'-UTR)) containing miR-132-binding site I and the miR-132-binding site II, were amplified by PCR and ligated into pGEM-T vector (Promega), and, subsequently, recombinant T-plasmids were digested with Hind III and Spe I and cloned into the 3' end of the pMIR-REPORT Luciferase vector with CMV promoter (Applied Biosystems) at Hind III and Spe I sites. These two plasmids were designed as pMIR-*StAR*-3UTR-132-I and pMIR-*StAR*-3UTR-132-II. Mutagenesis of the miR-132-binding sites was carried out according to the instructions supplied with the Quick-change II site-directed mutagenesis kit (Stratagene) employing recombinant T-plasmids containing *mStAR* 3'-UTR fragments. The mutated *mStAR* 3'-UTR fragments were digested with Spe I and Hind III

Table 1 Primers used for PCR gene cloning, site-directed mutagenesis and quantitative real-time PCR.

Plasmid/gene	Forward (F) and reverse (R) primers	Restriction enzyme sites
pcDNA6-m <i>Star</i> -V5	5'-CTGAAAGCTTATGTTCTCGCTACGTTCAAG-3'-F 5'-CTGATCTAGAACACTGGGCTCAGAGGCAG-3'-R	HindIII/XbaI
pMIR- <i>Star</i> -3UTR-132-I	5'-CTGAACTAGTGAATCACAGCTTCAGCAG-3'-F 5'-CTGAAAGCTTGCCTCTGCCACTTGAGTG-3'-R	HindIII/Spe I
pMIR- <i>Star</i> -3UTR-132-II	5'-CTGAACTAGTACAGCATCTGGCAGGAAG-3'-F 5'-CTGAAAGCTTAGCTCTACTGACTTCTAGC-3'-R	HindIII/Spe I
pMIR- <i>Star</i> -3UTR-132-IM	5'-TGTCATTTGACACATTAGATTCTTCTAAAAAGG-3'-F 5'-GCCTCTAAATACAGTAACTGTGTAATCTAAGAAA-3'-R	Mutagenesis
pMIR- <i>Star</i> -3UTR-132-IIM	5'-GATAAAGTGACCGTTCTATTGTTTATAG-3'-F 5'-CTATTTCACTGGCAAGATAACAAATATC-3'-R	Mutagenesis
Mouse DAX-1 (<i>Nr0b1</i>)	5'-AAGGGACCGTGCTCTTTAAC-3'-F 5'-GGATGTGCTCAGTAAGGATCTG-3'-R	qRT-PCR primers
Mouse SF-1 (<i>Nr5a1</i>)	5'-GTCTCAAGTTCCTCATCCTCTTC-3'-F 5'-GGGTAGTGACACAAGGTGTAAT-3'-R	qRT-PCR primers
Mouse AArhgap32 (<i>Arhgap32</i>)	5'-TTGAAGTGCCCGAGTTCTT-3'-F 5'-TATATCCCATCCACAATGCCATAC-3'-R	qRT-PCR primers
Mouse MeCP2 (<i>Mecp2</i>)	5'-ATGAGCCACTACAACCTTCAG-3'-F 5'-AGCTTTCTGTGTTCAACCTTC-3'-R	qRT-PCR primers
Mouse CtBP1 (<i>Ctbp1</i>)	5'-GGCAGCGGGTTTGACAATA-3'-F 5'-GCACACTGCGATGCCTAGATC-3'-R	qRT-PCR primers
Mouse HDAC3 (<i>Hdac3</i>)	5'-TCCCCTCATTTGTTCTCTG-3'-F 5'-TCCTTGCCACTGTACTTCATC-3'-R	qRT-PCR primers
Mouse StAR (<i>Star</i>)	5'-CGGAGCAGAGTGGTGTATC-3'-F 5'-TGAGTTTAGTTAGTCTGGAGGGACTTC-3'-R	qRT-PCR primers
Mouse 3β-HSD (<i>Hsd3b</i>)	5'-AGCGAGATCTGGGCTATGA-3'-F 5'-CCCTGTGCTGTTCCACTATT-3'-R	qRT-PCR primers
Mouse CYP11A1 (<i>Cyp11a1</i> ; P450 _{scc})	5'-CGCATCAAGCAGCAAAATTC-3'-F 5'-ATGCGCTCCCAATATAAC-3'-R	qRT-PCR primers
Mouse 20α-HSD (<i>Hsd17b1</i>)	5'-GCCTAAAGATGAGCATGGAAAC-3'-F 5'-TTAAAGTTAGACACCCGATGG-3'-R	qRT-PCR primers
Mouse 36B4 (<i>Rplp0</i>)	5'-GAAAAGGTCAAGGCCTTCCT-3'-F 5'-CGAAGAGACCGAATCCCAT-3'-R	qRT-PCR primers

Restriction enzyme sites are underlined. Gene symbols are listed in italics.

Arhgap32, rho GTPase activating protein 32; *Ctbp1*, C-terminal binding protein 1; *Hdac3*, histone deacetylase 3; *Hsd17b1*, hydroxysteroid 17-beta dehydrogenase 1; *Hsd3b*, hydroxy-delta-5-steroid dehydrogenase, 3beta-and steroid delta isomerase; *Mecp2*, methyl-CpG binding protein 2; *Nr0b1*, nuclear receptor subfamily 0 Group B member 1; *Nr5a1*, nuclear receptor subfamily 5 group A member 1; *Rplp0*, acidic ribosomal phosphoprotein P0; *Star*, steroidogenic acute regulatory protein.

and cloned into the 3' end of the pMIR-REPORT Luciferase vector. The mutated fragment plasmids were named as pMIR-StAR-3UTR-132-IM and pMIR-StAR-3UTR-132-IIM. The sequence of the inserted fragments was confirmed by sequencing. Primers for plasmid construction and mutagenesis are listed in [Table 1](#).

qRT-PCR analyses of miR-132 and miR-212

Total RNAs were extracted from adrenal tissues, cultured rat ovarian granulosa cells, MLTC-1 Leydig cells or Y-1 adrenal cells using miRNeasy Isolation Mini Kit (Qiagen) following the manufacturer's protocol. RNA samples were reverse transcribed using the mercury LNA universal RT microRNA PCR kit (Exiqon Inc., Woburn, MA, USA). Following cDNA synthesis, quantitative real-time PCR was performed on an ABI Prism 7900 HT system using SYBR Green Real-time Master Mix (Invitrogen) and microRNA LNA PCR primer sets for miR-132 and miR-212 (Exiqon Inc.). Fold-changes of mRNA levels were determined after normalization to internal control, *U6*mRNA levels, a small nuclear gene, using the $2^{-\Delta\Delta C_t}$ method ([Hu et al. 2013b](#)).

Quantitative RT-PCR (qRT-PCR)

Total RNA from Y-1 cells transfected with $\pm$ miR-132 mimic or scrambled oligonucleotide and treated with $\pm$ Bt₂cAMP was extracted using a miRNeasy Mini Kit (Qiagen). To measure mRNA expression, total RNA (2 μ g) from Y-1 cell samples was reverse transcribed using Superscript Reverse Transcriptase (Invitrogen) and oligo (dT) primers. The resulting complementary cDNA samples were used for qRT-PCR using SYBR Green Real-time Master Mix (Invitrogen) and specific primers as listed in [Table 1](#). qPCR was performed on an ABI Prism 7900 HT quantitative PCR system (Applied Biosystems). Target mRNA expression in each sample was normalized to the 36B4 signal. The $2^{-\Delta\Delta C_t}$ method was used to calculate relative mRNA expression levels.

Luciferase reporter assays

For the luciferase assay, CHO cells at a density of 1×10^5 per well in 24-well plates were co-transfected with pMIR-REPORT reporter constructs containing the 3' UTR of the StAR gene containing native or mutant site I- or site II-binding site for miR-132 $\pm$ miR-132 mimic or scrambled oligonucleotide using Lipofectamine 2000 transfection reagent (Invitrogen). Twenty-four hours after transfection,

cells were harvested and lysed with passive lysis buffer (Promega) according to the manufacturer's instructions. Reporter activity was measured on a SpectraMaxL luminescence microplate reader (Molecular Devices). The pRL-TK vector (Promega) *Renilla* luciferase control reporter vector, which provides constitutive expression of *Renilla* luciferase, was used as an internal control. The results are expressed as relative luciferase activities (firefly luciferase/*Renilla* luciferase).

Cell transfection

miR-132 mimic was purchased from Ambion. For transient transfections, Y-1 cells were plated 12 h before transfection at 50% confluence and then transfected with miR-132 mimic or scrambled oligonucleotide using Lipofectamine 2000 transfection reagent. For co-transfection experiments with miRNA and plasmid, pcDNA6-LacZ was used as control plasmid. Thirty-six to forty-eight hours after transfection, the cells were treated with $\pm$ Bt₂cAMP (2.5 mM) for 6 h and subsequently, the expression of DAX-1, SF-1, Arhgap32, MeCP2, CtBP1, HDAC3, StAR, 3 β -HSD, CYP11A1 and 20 α -HSD was assessed by qRT-PCR and/or Western blotting.

Western blot analyses

Y1 cells were harvested and homogenized in RIPA buffer (25mM Tris-HCl, pH 7.6, 150mM NaCl, 1% NP-40, 1% sodium deoxycholate and 0.1% SDS) supplemented with Thermo Scientific HaltProtease Inhibitor Cocktail (1mM AEBSE, 800nM aprotinin, 50nM bestatin, 15nM E-64, 5mM EDTA, 20nM leupeptin, 10nM pepstatin). The lysates were incubated for 20 min on ice and subsequently centrifuged at 10,000g for 5 min. Supernatants were collected, and protein concentrations were determined using the Pierce BCA Protein Assay Kit. Suitable aliquots of samples were mixed with equal volumes of 5 $\times$ Laemmli sample buffer (120mM Tris-HCl (pH 6.8), 2% SDS (wt/vol), 10% sucrose (wt/vol) and 1% 2-mercaptoethanol), and 10–20 μ g of total protein of each sample was subjected to 10% SDS-PAGE under denaturing conditions. Protein markers were also loaded on the gels. Following electrophoretic separation, the proteins were transferred to Immobilon-P PVDF Transfer membrane (EMD Millipore) using standard techniques. Blotted membranes were blocked with Odyssey blocking buffer (LI-COR Biosciences, Lincoln, NE, USA) for 1 h at room temperature and then incubated with anti-MeCP2, anti-SF-1, anti-StAR

or anti-GAPDH. After three washes with Tris-buffered saline containing 0.1% Tween 20, the membranes were incubated with IRDye 800CW goat anti-rabbit secondary antibody (LI-COR) for 1 h. Protein bands were detected with the Odyssey Infrared Imaging system (LI-COR Biosciences).

Measurement of progesterone and 20 α -OHP secretion

Y1 cells were transfected with $\pm$ miR-132 mimic or scrambled oligonucleotide for 48 h and were subsequently treated with $\pm$ Bt₂cAMP (2.5 mM) or 22(R)-hydroxycholesterol (10 μ M) for 6 h. After incubation, progesterone and 20 α -OHP in the medium was determined by radioimmunoassay (Azhar *et al.* 1998).

Thin-layer chromatography

Y1 cells were transfected with $\pm$ miR-132 mimic or scrambled oligonucleotide for 48 h and were subsequently incubated with [³H] pregnenolone (1 μ Ci/mL) + unlabeled pregnenolone (10 μ M) for 6 h. After incubation, media and cell samples were extracted with diethyl ether, and the ether phase in each case after drying under nitrogen was subjected to thin-layer chromatography using chloroform-ether (5:1, v/v) as described by Jones and Hsueh (Jones & Hsueh 1982). The [³H] progesterone and [³H] 20 α -OHP spots were scraped from the plate and counted for the determination of radioactivity.

Statistical analysis

The results are presented as the means $\pm$ S.E. for at least three independent experiments. All data were tested for

normality (Kolmogorov–Smirnov test) and homogeneity of variances (Bartlett's test). We used one-way ANOVA or two-way ANOVA to analyze the corresponding data when the assumptions for parametric analyses were met. Statistical analyses were performed using ANOVA followed by the Bonferroni's post-test using GraphPad Prism Software, Prism 6 (GraphPad Software). Descriptive statistics are presented as mean $\pm$ standard error, and the significance level is set at $\alpha=0.05$.

Results

Relative expression and hormonal regulation of miR-132 and miR-212

miR-132 and miR-212 are derived from the same primary microRNA transcript and these two miRNAs share the same seed sequence (Wanet *et al.* 2012, Remenyi *et al.* 2013). We first determined the relative expression of these two microRNAs by qRT-PCR under basal conditions and in response to hormonal stimulation of rat adrenals, Y1 mouse adrenocortical cells, MLTC-1 mouse Leydig tumor cells and primary rat ovarian granulosa cells. Dishes of MLTC-1 and Y1 cells treated with or without Bt₂cAMP (2.5 mM) for 6 h, and granulosa cells treated with or without Bt₂cAMP (2.5 mM) for 24 h, were subjected to RNA isolation followed by qRT-PCR. Likewise, groups of rats were treated subcutaneously with ACTH (10 IU daily for 4 days) or a vehicle (control), and adrenal samples were subjected to RNA isolation and qRT-PCR. As shown in Fig. 1, miR-132 was expressed at much higher basal levels than miR-212 in MLTC-1, Y1 and granulosa cells and the adrenal gland (approximately 300-, 250-, 270- and 220-fold, respectively). Treatment of MLTC-1, Y1 and

Figure 1

Basal expression and hormonal regulation of miR-132 and miR-212 in mouse MLTC-1 Leydig tumor cells, rat ovarian granulosa cells, mouse Y1 adrenal cells and rat adrenal glands. Cultured MLTC-1 and Y1 cells were treated with $\pm$ Bt₂cAMP (2.5 mM) for 6 h and granulosa cells $\pm$ Bt₂cAMP (2.5 mM) for 24 h, and isolated RNA samples analyzed for miR-132 and miR-212 expression by qRT-PCR. Likewise, groups of rats were treated subcutaneously with ACTH (10 IU every 24 h for 4 days) or a vehicle (control), and adrenal samples were subjected to total RNA isolation. Isolated RNA samples were analyzed for expression of miR-132 and miR-212 by qRT-PCR. miR-132 and miR-212 were normalized to the housekeeping gene U6. We used one-way ANOVA to compare the differences between control and treated samples. Data are presented as means $\pm$ S.E. ($n=4$). * $P<0.05$; ** $P<0.01$; *** $P<0.001$.

granulosa cells with Bt_2cAMP or adrenals with ACTH *in vivo* upregulated the expression of both miRNAs, although the extent of stimulation varied between approximately 0.4-fold to 5-fold (Fig. 1). In view of the very low expression of miR-212, we focused further studies on miR-132.

Increased miR-132 expression was associated with repression of StAR protein expression, activation of steroidogenic enzymes and modulation of steroidogenesis in Y1 cells

The above results showing that miR-132 expression is increased in intact adrenals and the steroidogenic cells, granulosa, MLTC-1 and Y1 cells and is increased in response to trophic hormone or Bt_2cAMP (Fig. 1) raised the possibility that miR-132 is involved in the regulation of steroidogenesis. To test this possibility, we transfected Y1 cells with miR-132 mimic or scrambled oligonucleotide, and cells were subsequently analyzed for the expression of putative miR-132 target genes (Arhgap32, MeCP2, CtBP1 and HDAC3), steroidogenic transcription factors (DAX-1 and SF-1), StAR protein (9,10,14,19), steroidogenic enzymes (CYP11A1, 3 β -HSD and 20 α -HSD) and steroid (progesterone and 20 α -OHP) secretion both under basal conditions and in response to cAMP stimulation. qRT-PCR results demonstrated that overexpression of

miR-132 under basal conditions increased mRNA levels of 3 β -HSD and 20 α -HSD in Y1 cells. The enzyme 3 β -HSD catalyzes the conversion of pregnenolone to progesterone, whereas 20 α -HSD catalyzes the conversion of progesterone into its inactive form 20 α -OHP. However, no changes in mRNA levels were noted for DAX-1, SF-1, Arhgap32, MeCP2, CtBP1, HDAC3, StAR and CYP11A1 (Fig. 2A). In contrast, Western blot analysis showed reduced protein levels of MeCP2 and StAR in cells transfected with miR-132 (Fig. 2B). To investigate the effects of miR-132 on steroid production, we collected media from the cultured miR-132 and scrambled oligonucleotide-transfected cells and assessed progesterone and 20 α -OHP concentrations (Fig. 2C). Overexpression of miR-132 significantly decreased the basal levels of both secreted progesterone and 20 α -OHP.

We next assessed the effects of cAMP treatment of control (scrambled oligonucleotide) and miR-132 transfected Y1 cells on the mRNA levels of various proteins noted above and on the secreted levels of two progestins, progesterone and 20 α -OHP. Like basal transfected cells, miR-132-transfected and Bt_2cAMP -treated cells also showed increased mRNA levels of 3 β -HSD and 20 α -HSD, while mRNA levels of other proteins were not affected (Fig. 2A). Likewise, overexpression of miR-132 decreased MeCP2 and StAR protein levels in cAMP-stimulated cells,

Figure 2

Effects of miR-132 overexpression on putative miR-132 sensitive genes and steroid production in Y1 cells. Y-1 adrenal cells were plated 12 h before transfection at 50% confluence and then transfected with miR-132 mimic or scrambled oligonucleotide. Thirty-six to 48 h after transfection, the cells were incubated under basal or cAMP-stimulated (2.5 mM) conditions for 6 h and, subsequently, the expression of DAX-1, SF-1, Arhgap32, MeCP2, CtBP1, HDAC3, StAR, 3 β -HSD, CYP11A1 and 20 α -HSD measured by qRT-PCR (A) and/or Western blotting (B) as described under the Experimental Section. The media samples without (C) or with (D) cAMP treatment were analyzed for progesterone and 20 α -hydroxyprogesterone (20 α -OHP) content by radioimmunoassay. We used two-way ANOVA to compare the differences among different samples. Data are presented as means $\pm$ s.e. ($n=4$). * $P<0.05$; ** $P<0.01$; *** $P<0.001$.

although cAMP treatment rescued the miR-132 reduced StAR protein to the basal level (Fig. 2B). To investigate the function of miR-132-induced increases in β HSD and 20 α -HSD activity, we analyzed the secreted levels of progesterone and 20 α -OHP. Compared with control (scrambled oligonucleotide), the overexpression of miR-132 increased the levels of both progesterone and 20 α -OHP in the culture medium in response to cAMP treatment (Fig. 2D). Furthermore, the cells secreted roughly 25-fold more 20 α -OHP into the medium relative to progesterone. These results demonstrate that miR-132 overexpression promotes the formation of biologically inactive progesterone, 20 α -OHP, by enhancing the functional expression of 20 α -HSD activity.

To further study miR-132 stimulation of 20 α -OHP production, we followed the sequential conversion of [3 H] pregnenolone to [3 H] progesterone and [3 H] 20 α -OHP and the formation of progesterone and 20 α -OHP from 22(R)-hydroxycholesterol in Y1 cells transfected with miR-132 (or control scrambled oligonucleotide). 22(R)-hydroxycholesterol is a freely diffusible cholesterol derivative, which does not require StAR protein for its transport to mitochondria for catabolism by the P450_{sc} (CYP11A) enzyme for the formation of the common steroid precursor pregnenolone. Results presented in Fig. 3 demonstrate that transfection of Y1 cells with

Figure 3

Effects of miR-132 on cellular conversion of [3 H] labeled pregnenolone and 22(R)-hydroxycholesterol to progestins in Y1 cells. Y-1 adrenal cells were plated 12 h before transfection at 50% confluence and then transfected with miR-132 mimic or scrambled oligonucleotide. Thirty-six to 48 h after transfection, the cells were incubated with [3 H] pregnenolone (1 μ Ci/mL; 10 μ M) or 22(R)-hydroxycholesterol (10 μ M) for 6 h, the precursor and free diffusible cholesterol substrate, respectively. (A), [3 H] pregnenolone-treated media samples were quantified for the accumulated levels of [3 H] progesterone/[3 H] 20 α -OHP by thin-layer chromatography (TLC). (B) 22(R)-hydroxycholesterol treated samples were analyzed for progesterone and 20 α -OHP by radioimmunoassay (RIA) techniques, respectively. For these studies, transfected cells were not further stimulated with cAMP. One-way ANOVA was used to assess the differences between control and miR-132 transfected samples. Data are presented as means $\pm$ s.e. (n=4). *P<0.05.

miR-132 increased the conversion of [3 H] pregnenolone into [3 H] 20 α -OHP (Fig. 3A) and preferentially enhanced the production of 20 α -OHP from 22(R)-hydroxycholesterol (Fig. 3B). These data further confirm the finding that miR-132 promotes the production of 20 α -OHP by upregulating the functional expression of 20 α -HSD.

miR-132 suppresses StAR protein expression by targeting specific sites within the 3' UTR of StAR

The results presented in Fig. 2A and 2B raised the possibility that miR-132 suppression of basal steroidogenesis might be linked to the observed inhibition of StAR protein expression (Fig. 2B). Here, we directly assessed the binding of the miR-132 to the 3'-UTR of StAR and regulation of its expression level by carrying out a luciferase reporter assay. Individual fragments of the 3' UTR region of the StAR gene containing site I, mutated site I, site II or mutated site II were inserted downstream of the luciferase open reading frame of pMIR-REPORT vector (Fig. 4A). CHO cells were co-transfected individually with StAR-3'-UTR (site I, site II or respective mutated sites) $\pm$ miR-132 for 36 h followed by determination of luciferase activities. Overexpression of

A mmu-miR-132-3P

Mice StAR 3'UTR Site I (WT) 5' -AGATTATGTCATTGACTGTTT-3'
StAR 3'UTR Site I (MUT) 5' -AGATTATGTCATTGACACATT-3'
mmu-miR-132-3p Site I 3' -GCUGGUACCGACAUUCUGACAAU-5'
Mice StAR 3'UTR Site II (WT) 5' -ATGTTAAGATAAAGTGACTGTTC-3'
StAR 3'UTR Site II (MUT) 5' -ATGTTAAGATAAAGTGACCGTTC-3'
mmu-miR-132-3p Site II 3' -GCUGGUACCGACAUUCUGACAAU-5'

Figure 4

miR-132-binding site I in the 3' UTR of the mouse StAR gene mediates the downregulation of StAR expression by miR-132. (A) Seed sequences of the putative miR-132-binding sites (site I and site II) in the 3' UTR of mouse StAR. For the reporter gene assay, the 3' UTR region of the StAR gene-containing site I, mutated site I, site II or mutated site II for miR-132 was inserted downstream of the luciferase open reading frame of pMIR-REPORT vector. (B) CHO cells were co-transfected with the StAR 3' UTR (containing putative site I or site II or respective mutated sites for miR-132 binding) $\pm$ miR-132 mimic. Reporter assays were performed using a dual luciferase kit. Two-way ANOVA was used to compare the differences among different samples. The results are expressed as mean relative luciferase activities $\pm$ s.e. (n=4). *P<0.05.

Figure 5

Effects overexpression of StAR and miR-132 in Y1 cells on basal progestin production. Y1 cells were transiently transfected with 50 nM scrambled oligonucleotide or miR-132 mimic plus mouse StAR-V5 cDNA construct. After 48 h of transfection, medium was changed to basal medium, dishes incubated for an additional 6 h and subsequently medium samples collected and analyzed for progesterone and 20α-OHP by radioimmunoassay. Cell samples were analyzed for overexpressed StAR protein levels by Western blotting. StAR protein levels were normalized to GAPDH. (A) StAR protein expression; (B) secreted progesterone and (C) secreted 20α-OHP. We used one-way ANOVA to compare the differences between control and miR-132 co-transfected samples. Data are presented as means ± s.e. (n=4). *P<0.05.

miR-132 in CHO cells significantly decreased the luciferase activity (Fig. 4B). This inhibition was abolished when binding site I was mutated. However, no inhibitory effect of miR-132 on the activity of the 3' UTR of StAR reporter construct containing site II was detected, indicating that site I is the target of miR-132 action on StAR 3' UTR (Fig. 4B). The results presented in Fig. 5 demonstrate that overexpression of StAR/StarD1 overcomes the inhibitory action of miR-132 on basal progestin production, and, in fact, increased 20α-OHP production.

miR-132 upregulates the expression of 3β-HSD and 20α-HSD activities and progestin production by inhibiting MeCP2 expression

Next, we investigated the potential mechanism by which miR-132 upregulates the expression of 3β-HSD and 20α-HSD activities. Since MeCP2, a transcriptional regulatory protein (Hite *et al.* 2009), was previously shown to be a downstream target of miR-132 in the brain (Wanet *et al.* 2012) and our current observation that overexpression of miR-132 expression suppressed MeCP2 protein expression in Y1 cells (Fig. 2B), this raised the possibility that this transcription factor plays a regulatory role in miR-132 induction of 3β-HSD and 20α-HSD enzymes. Initially, we evaluated the separate and combined effects of transfection of Y1 cells with miR-132 and MeCP2 constructs on the mRNA expression of 3β-HSD and 20α-HSD. Quantitative RT-PCR data presented in Fig. 6 (A and B) show that overexpression of miR-132 increased mRNA levels of both 3β-HSD and 20α-HSD, but, as before, had no effect on mRNA levels of other genes examined both under basal conditions and in response to cAMP stimulation. In contrast, simultaneous expression

of MeCP2 reversed the increased effect on mRNA levels of 20α-HSD in control and cAMP-stimulated cells. Western blotting data presented in Fig. 6C confirmed the miR-132-induced suppression of MeCP2 protein expression and the increased expression of MeCP2 protein in cell extracts from cells transfected with the MeCP2 cDNA construct. Likewise, overexpression of miR-132 alone or miR-132 plus MeCP2 decreased progesterone production under basal conditions, but increased cAMP-stimulated progesterone secretion (Fig. 6D). The production of 20α-OHP, however, was significantly decreased in cells co-transfected with miR-132 along with MeCP2 as compared to miR-132 alone (Fig. 6E). These studies lead us to conclude that miR-132 induces the expression of two pregnenolone metabolizing enzymes, particularly 20α-HSD, through inhibition of MeCP2 protein expression.

To further examine the contribution of MeCP2 protein in miR-132-mediated induction of 3β-HSD and 20α-HSD activities, we employed a siRNA strategy to deplete mRNA and protein levels of MeCP2 in Y1 cells. Quantitative RT-PCR results demonstrated that transfection of Y1 cells with MeCP2-specific siRNA reduced MeCP2 mRNA levels by 60–70% (Fig. 7A). Western blot analysis showed that Y1 cells had approximately 50% MeCP2 protein levels when transfected with MeCP2 siRNA (Fig. 7B). In contrast, mRNA levels of 3β-HSD and 20α-HSD were significantly increased (more robustly for 20α-HSD) following transfection of Y1 cells with MeCP2 siRNA (Fig. 7A). We also investigated the effect of siRNA-induced depletion of MeCP2 on steroid (progesterone and 20α-OHP) production. Compared to negative control, transfection of Y1 cells with MeCP2 siRNA increased the basal production of 20α-OHP (Fig. 7D). In contrast, no such stimulatory effect was noted on progesterone production (Fig. 7C). These studies

Figure 6

Effects of miR-132 and MeCP2 overexpression on StAR, CYP11A1, 3 β -HSD and 20 α -HSD expression and progesterin production. Y1 cells were transiently transfected with scrambled oligonucleotide or miR-132 mimic $\pm$ mouse MeCP2-V5 cDNA construct or control plasmid pcDNA6-LacZ. Thirty-six to 48 h after transfection, the cells were incubated in fresh medium for 6 h without or with cAMP. The mRNA expression of StAR, MeCP2, CYP11A1, 3 β -HSD and 20 α -HSD in basal (A) or cAMP treated (B) samples was measured by qRT-PCR. (C) Protein levels were analyzed by Western blotting as described under the Experimental Section. The media samples were analyzed for progesterone (D) and 20 α -OHP (E) content by radioimmunoassay. One-way ANOVA was used to analysis the differences among different samples as showed in the figures. Data are presented as means $\pm$ s.e. ($n=4$). * $P<0.05$; ** $P<0.01$.

led us to conclude that miR-132 upregulates 3 β -HSD and 20 α -HSD and increases 20 α -OHP by inhibiting MeCP2 protein expression. These studies also establish that MeCP2 is a negative regulator of the expression of 20 α -HSD and also possibly 3 β -HSD.

To further confirm the physiological significance of miR-132 regulation of steroidogenesis, we also carried out loss-of-function and gain-of-function studies using primary rat ovarian granulosa cells. We transfected granulosa cells with miR-132 mimic and anti-miR-132 for 6 h and subsequently cells were maintained in the basal medium for 24 h. Following addition of Bt2cAMP $\pm$ hHDL3 for 24 h, the medium samples were analyzed for 20 α -OHP content by specific RIAs. As shown in Fig. 8, overexpression of miR-132 increased 20 α -OHP production with and without HDL loading, while knockdown of miR-132 resulted in a significant decrease of 20 α -OHP production from the granulosa cells.

Discussion

In these studies, we evaluated the role of trophic hormone regulated miR-132 in the regulation of steroidogenesis using a model Y1 adrenal cell line. Although miR-132 together with miR-212 comprises the evolutionary conserved miR-132/212 family, miR-212 expression was almost negligible compared to miR-132 not only in Y1 cells, but also in rat ovarian granulosa cells, mouse Leydig tumor MLTC-1 cells and the rat adrenal gland. Thus, our studies focused on miR-132. The results demonstrated that miR-132 itself is positively regulated

Figure 7

Effects of MeCP2 siRNA on MeCP2, StAR, CYP11A1, 3 β -HSD, and 20 α -HSD expression and progesterin production in Y1 cells. Y1 cells were transiently transfected with mouse MeCP2 siRNA or control (scrambled) siRNA. Thirty-six to 48 h after transfection, the cells were incubated in fresh medium for 6 h and, subsequently, the expression of StAR, MeCP2, CYP11A1, 3 β -HSD and 20 α -HSD was measured by qRT-PCR (A) and/or Western blotting (B) as described under Experimental Section. The media samples were analyzed for progesterone (C) and 20 α -OHP (D) content by radioimmunoassay. We used one-way ANOVA to compare the differences between control and siMeCP2 transfected samples. Data are presented as means $\pm$ s.e. ($n=4$). * $P<0.05$; ** $P<0.01$, *** $P<0.001$.

Figure 8

Effect of treatment of rat ovarian granulosa cells with mimic or anti-miR-132 on 20α-OHP production. Primary rat granulosa cells were transfected with ± miR-132 mimic, anti-miR-132 or scrambled oligonucleotide using Polyjet™ for 6 h, and cells were maintained in basal medium for 24 h before treatment with Bt2cAMP (2.5 mM) ± HDL (500 μg/mL) for an additional 24 h. At the end of incubation, medium samples were analyzed for 20α-OHP content by radioimmunoassay. We used one-way ANOVA to compare the differences between control and treated samples. Data are presented as means ± s.e. ($n=4$). * $P<0.05$; ** $P<0.01$, *** $P<0.001$.

by cAMP, the second messenger of trophic hormone, in Y1, granulosa and MLTC-1 cells and also by ACTH in the rat adrenal gland *in vivo*. Studies revealed that miR-132 inhibits steroidogenesis by a dual mechanism. Under basal conditions, miR-132 attenuates steroidogenesis by reducing StAR protein levels. In response to hormonal stimulation, miR-132 induces the expression of 3β-HSD and 20α-HSD through inhibition of transcription factor MeCP2. This, in turn, causes sequential conversion of the CYP11A1 reaction product pregnenolone to progesterone and then biologically inactive 20α-OHP (Fig. 9). Together, these data indicate that miR-132 plays a key role in the posttranscriptional regulation of steroid hormone biosynthesis.

miR-132 expression is transcriptionally induced by cAMP-responsive element-binding (CREB) protein (Impey *et al.* 2004, Vo *et al.* 2005, Wanet *et al.* 2012) and inhibited by repressor element 1 silencing transcription factor/neuron-restrictive silencer factor (REST/NRSF) (Conaco *et al.* 2006, Wanet *et al.* 2012). miR-132 has been most extensively studied in the brain, where it is implicated in neuronal differentiation and function (Impey *et al.* 2004, Vo *et al.* 2005, Wu & Xie 2006, Conaco *et al.* 2006, Cheng *et al.* 2007, Wayman *et al.* 2008). Increasing evidence now suggests that miR-132 also regulates multiple processes in non-neuronal tissues. For example, miR-132 expression is induced by lipopolysaccharide (LPS)

Figure 9

Sites of action of miR-132 on the steroidogenic pathway. Based on the data obtained we propose that miR-132 regulates steroidogenesis by: (1) inhibiting StAR protein expression and StAR-mediated cholesterol transport to mitochondria and (2) suppressing MeCP2 expression to cause primarily increased expression of 20α-HSD and consequently 20α-HSD catalyzed increased conversion of progesterone into an inactive metabolite, 20α-OHP. Also, we believe that the first pathway is more physiologically relevant.

in THP-1 cells (Taganov *et al.* 2006) and is overexpressed in chronic lymphoblastic leukemia (Calin *et al.* 2004). More recent studies suggest that miR-132 regulates innate immunity (Lagos *et al.* 2010), regulates nutritional stress-induced chemokine production (Strum *et al.* 2009) and it (along with family member miR-212) contributes to the pathogenesis of cardiac hypertrophy and cardiomyocyte autophagy (Ucar *et al.* 2012). We found that the expression of miR-132 was upregulated in Y1 adrenal cells, ovarian granulosa cells and testicular Leydig tumor MLTC-1 cells in response to second messenger, cAMP, although the extent of cAMP stimulation of miR-132 expression varied considerably among cell types, as well as in adrenals of rats treated *in vivo* with ACTH. cAMP-mediated modulation of miR-132 expression has also been reported in non-steroidogenic cells; the cAMP analog 8-Br-cAMP-induced miR-132 expression in primary lymphatic endothelial cells (LECs), whereas the cAMP antagonist RpAMPs partially blocked the induction of Kaposi's sarcoma-associated herpes virus (KSHV)-infected LECs (Lagos *et al.* 2010).

Although miR-132 and miR-212 comprise the evolutionary conserved miR-132/212 cluster (Wanet *et al.* 2012, Remenyi *et al.* 2013), and these microRNAs share

the same transcript and the same 5' seed sequence in humans and rodents, qRT-PCR measurements indicated that miR-212 was expressed several hundredfold lower than miR-132 both under basal conditions and in response to cAMP stimulation. The data are consistent with other reports where only miR-132 was found to be induced and functional in various biological systems (Calin *et al.* 2004, Taganov *et al.* 2006, Cheng *et al.* 2007, Klein *et al.* 2007). However, a previous report provided evidence that both miR-132 and miR-212 were highly upregulated in ovulatory mouse mural granulosa cells *in vivo* in response to hCG treatment and under an *in vitro* culture condition following cAMP stimulation (Fiedler *et al.* 2008). At present, we are unable to provide an exact explanation for the differences between our results using rat granulosa cells with this report (Fiedler *et al.* 2008) that employed mouse mural granulosa cells. It may be related to species differences, granulosa cell types or both.

Our studies demonstrate that miR-132 negatively regulates steroidogenesis in Y1 cells by a novel dual mechanism. miR-132 can directly repress StAR protein expression through its binding to site I (GACTGTT), but not site II, in the 3' UTR of the mouse *StAR* gene, thereby negatively regulating StAR-mediated cholesterol transport to mitochondria for cholesterol side-chain cleavage catalyzed by CYP11A1 for pregnenolone production and, consequently, progesterone (steroid) production. We further demonstrated that overexpression of miR-132 reduced the protein levels of StAR protein without impacting its mRNA levels both under basal conditions and in response to cAMP stimulation. These results, thus, validate StAR as a direct target of miR-132. The observed reduction in StAR protein levels is possibly due to miR-132 interference with the translation process. At a functional level, the inhibition of basal levels of StAR protein is reflected by corresponding decreases in the basal production of progesterone and 20 α -OHP in Y1 cells transfected with miR-132.

Our studies also indicate that treatment of cells overexpressing miR-132 with cAMP increased the secretion of stimulated progesterone and 20 α -OHP secretion, as a result of increased expression of 3 β -HSD and 20 α -HSD, and cAMP-stimulated StAR protein expression. Although in the current studies we have identified StAR protein as a downstream target of miR-132, it appears that cAMP-stimulated cells, despite significant inhibition in cAMP-induced StAR protein levels by miR-132, still maintain more than an adequate amount of this protein to maximally stimulate steroidogenesis.

This, coupled with increased expression of 3 β -HSD and 20 α -HSD, assures a rapid 3 β -HSD-catalyzed conversion of CYP11A1 reaction product pregnenolone to biologically active progesterone and, subsequently, quantitative conversion of progesterone to inactive 20 α -OHP, as a result of the miR-132-mediated robust induction of 20 α -HSD. Additional studies confirmed these changes by demonstrating that transfection of Y1 cells with miR-132 increased the conversion of [³H] pregnenolone into [³H] 20 α -OHP and preferentially enhanced the production of 20 α -OHP from 22(R)-hydroxycholesterol, further confirming that miR-132 promotes the production of 20 α -OHP by upregulating the functional expression of 3 β -HSD and 20 α -HSD enzymes. Because cultured Y1 cells, both under basal conditions and in response to trophic hormone stimulation, primarily synthesize and secrete 20 α -hydroxy- Δ 4-pregnene-3-one (20 α -OHP) and 11 β ,20 α -dihydroxy- Δ 4-pregnene-3-one (11 β , 20 α -OHP), along with small amounts of progesterone (Rainey *et al.* 2004), it was important to document that overexpression or knockdown of miR-132 altered 20 α -OHP production in cells that primarily secrete progesterone as their steroid product. Indeed, overexpression or knockdown of miR-132 in primary rat ovarian granulosa cells increased and decreased 20 α -OHP production, respectively, thus confirming that the observed effects on steroid production were not due to unique characteristics of Y1 cells.

In addition, we observed that overexpressed miR-132 inhibited MeCP2 protein expression, a well-established downstream target of miR-132 (Klein *et al.* 2007), which raised the possibility that MeCP2 might be involved in miR-132-mediated induction of 3 β -HSD and 20 α -HSD. This possibility was confirmed with the use of MeCP2-specific siRNA. Transfection of Y1 cells with MeCP2-specific siRNA reduced MeCP2 mRNA levels by 60–70% and protein levels by ~50%. Under these experimental conditions, the mRNA levels of 3 β -HSD and 20 α -HSD were significantly increased (more robustly for 20 α -HSD). Likewise, transfection of Y1 cells with MeCP2 siRNA increased the basal production of 20 α -OHP. These studies led us to conclude that miR-132 upregulates 3 β -HSD and 20 α -HSD mRNA expression and increases 20 α -OHP potentially by inhibiting MeCP2 protein expression. Furthermore, direct experimental evidence is needed to demonstrate whether MeCP2 directly inhibits 3 β -HSD and/or 20 α -HSD activity. These studies also establish that MeCP2 is a negative regulator of the expression of 3 β -HSD and 20 α -HSD enzymes.

In summary, these studies demonstrate for the first time that a trophic hormone-regulated microRNA, miR-132, regulates steroidogenesis by a novel dual mechanism. It attenuates steroidogenesis by repressing StAR protein expression and by causing the induction of 20 α -HSD via inhibition of MeCP2, which results in the generation of biologically inactive steroid. Our studies suggest that miR-132 interferes with steroidogenesis not only by upregulating 20 α -HSD, but also by limiting the availability of cholesterol precursor for side-chain cleavage via inhibition of StAR. Further studies are required to determine how tissue-specific trophic hormones, which positively regulate almost all processes linked to steroidogenesis, also control the events that are connected with the inhibition of the steroidogenic process. In addition, further studies will be needed to further explore the roles of miR-132 and minor miR-212 in trophic hormone regulation of adrenal and ovarian steroidogenesis and expression, steroid profiles and expression of 3 β -HSD and 20 α -HSD *in vivo*. Finally, the data suggest that under conditions of markedly elevated ACTH, the increased miR-132, and to a very lesser degree miR-211, expression tends to restrain the maximum glucocorticoid output by the adrenals, thus potentially leading to a suboptimal glucocorticoid response in some clinical settings.

Declaration of interest

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

Funding

This work was supported, in whole or in part, by National Institutes of Health, NHLBI, Grant, 2R01HL33881. This work was also supported by the Office of Research and Development, Medical Service, Department of Veterans Affairs.

Author contribution statement

S A, Z H, F K and W J S conceived and designed the experiments. Z H and W J S performed the experiments. Z H and W J S analyzed the data. Z H, S A, F K and W J S wrote the paper. Z H, W J S, F K and S A edited the manuscript.

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Received in final form 6 July 2017

Accepted 20 July 2017

Accepted preprint published online 20 July 2017