



A partial loss-of-function variant in *GNRHR* gene in a Chinese cohort with idiopathic hypogonadotropic hypogonadism

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Background: Idiopathic hypogonadotropic hypogonadism (IHH) is a rare genetic disease attributed to the disorder of hypothalamic-pituitary-gonadal axis. Mutations in the *GNRHR* gene are one of the most common genetic causes of IHH. Herein, we aimed to investigate *GNRHR* variants in a Chinese cohort with IHH, and to characterize them at the molecular level.

Methods: A total of 153 IHH patients were recruited, and variants were detected using a tailored next-generation sequencing panel. *GNRHR* rare sequencing variant (RSV) was verified using Sanger sequencing. Phenotypic features and therapeutic outcomes of patients were followed up. In order to examine the pathogenicity of the *GNRHR* RSV, we performed conservative analysis, crystal structure prediction, expression analysis as well as the assessment of ERK1/2 activation and IP3/Ca²⁺ response.

Results: The same heterozygous RSV (p.R240Q) in *GNRHR* was identified in four sporadic IHH patients. These patients exhibited different severity of testicular development and hormone profile. hCG treatment was effective in improving gonadal development, serum testosterone, and semen quality. The *GNRHR* RSV has no effect on the expression of mRNA and protein, whereas damaged ERK1/2 activation and inositol triphosphate/calcium signaling.

Conclusions: The study expands *GNRHR* mutation spectrum in IHH patients, and reveals that the *GNRHR* RSV is a partial loss-of-function mutation. Although this heterozygous RSV may not have a significant influence on the pathogenesis of IHH, but its homozygous/ compound status should be paid attention in this research field.

Keywords: Idiopathic hypogonadotropic hypogonadism (IHH); rare sequencing variant; *GNRHR*; heterozygote; phenotype

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Introduction

Patients with idiopathic hypogonadotropic hypogonadism (IHH) present with incomplete puberty development, low gonadotropin levels, and infertility. Most patients are

diagnosed in their late adolescence or in early adulthood. IHH is a rare genetically heterogeneous disorder caused by the deficient development and migration of gonadotropin-releasing hormone (GnRH) neurons and deficient GnRH

secretion and action (1). The impaired hypothalamic-pituitary-gonadal axis further leads to steroidogenic and gametogenic disorders. The prevalence of IHH is approximately 1:10,000 in males and 1:50,000 in females (2). IHH has high phenotypic and genetic heterogeneity. IHH with olfactory disorder (50–60% of IHH) is known as Kallmann syndrome (KS), while IHH with a normal sense of smell is described as normosmic IHH (nIHH) (3). There are multiple inheritance mode of causal genes, including autosomal dominant, autosomal recessive, X-linked recessive, as well as digenic and oligogenic transmission (4).

To date, approximately >30 genes has been discovered to be implicated in IHH, including *ANOS1*, *GNRH1/GNRHR*, *FGF8/FGFR1*, *PROK2/PROKR2*, *SEMA3A/PLXNA1*, *KISS1/KISS1R*, *SEMA3E*, *NSMF*, *HS6ST1*, *WDR11*, *SOX10*, etc. (5). *GNRHR* (GnRH receptor) is the first gene identified as a monogenic cause of nIHH (6). *GNRHR* is located in chromosome 4q21.2, and encodes a seven-transmembrane G protein coupled receptor (7). *GNRHR* and its ligand, GnRH, play a crucial role in the secretion of anterior pituitary gonadotropes (8). Their binding facilitates an increase in inositol trisphosphate and calcium mobilization, and ultimately triggers the synthesis and secretion of the follicle-stimulating and luteinizing hormones (9). So far, at least fifty-five variants in the *GNRHR* gene have been found among nIHH patients, and the homozygous and compound heterozygous mutations are very common (10). However, it is noteworthy that heterozygous *GNRHR* mutations were often identified among IHH patients (11,12). A genetic study on a large IHH cohort showed that frequency of heterozygous *GNRHR* RSVs reached 2.5% (22/863), significantly higher than the healthy control population (12). An recent study also revealed that this heterozygous RSV was found in the three sporadic IHH patients, without carrying other common IHH gene mutations (11). Therefore, researchers speculated that monoallelic *GNRHR* RSVs are associated with IHH, but there was limited literature regarding their functional implications.

The identification of the more *GNRHR* RSVs is not only beneficial to the early genetic diagnosis of IHH, but also conducive to the development of individualized treatment for patients. For example, if the IHH patients with pathogenic *GNRHR* RSVs receive pulsatile GnRH treatment, it is predicted that their therapeutic response would not be very good. Gonadotropin treatment will help these patients achieve better therapeutic results. In addition, GnRH agonists (GnRH-a) are widely used in the treatment of prostate cancer, endometriosis and uterus adenomyosis.

It is reasonable to speculate that the *GNRHR* RSVs have a negative effect on the response to GnRH-a in these diseases. The aim of this study was to investigate *GNRHR* mutation profile in a Chinese IHH cohort, and to characterize the clinical manifestation and therapeutic outcome. In vitro functional analysis of *GNRHR* rare sequencing variants (RSVs) was also performed. We present the following article in accordance with the MDAR checklist (available at <http://dx.doi.org/10.21037/tau-20-1390>).

Methods

Patients

From 2005 to 2013, a total of 153 male patients were recruited at the Urology Clinic of Tongji Hospital, Wuhan, China. IHH was diagnosed according to standard diagnostic criteria (1), and adult-onset IHH (AOIHH) was diagnosed based on previous published criteria (13). The study protocol (IRB ID: 20150302) (including collection of blood samples) was approved by the Ethics Committee of Huazhong University of Science and Technology in compliance with the Declaration of Helsinki (as revised in 2013). Informed consent was signed with each patient.

Panel sequencing and frequency analysis

Genomic DNA was extracted from peripheral blood lymphocytes in all patients. To focus on the genetic variations in IHH-associated genes, we designed a tailored next-generation panel which included 31 causal genes and 52 candidate genes, as described in our previous study (14). The detected gene loci included exon and proximal intron DNA sequences ≤ 15 bp from splice site. Non-synonymous rare sequencing variants (RSVs) with minor allele frequency <1% (ESP6500, 1000genome, gnomAD databases) were included in our study. Sanger sequencing was performed to confirm all variants by using an ABI PRISM 3130 genetic analyzer (Applied Biosystems, Foster City, CA, USA). The primers were as follows: *GNRHR*-forward: 5'-GACAGCTCTGGACAGACAAA-3'; *GNRHR*-reverse: 5'-ACACTAACTCTAAGGAATACATACCG-3'. Variants were reported in agreement with Human Genome Variation Society nomenclature (15).

Clinical evaluation

At first hospital visit, we performed detailed phenotyping

and collected the medical history for each patient. Olfactory function was assessed by the olfactory testing or self-reporting. Tanner stage was evaluated by a senior physician according to Five Tanner stages of pubertal development (16). We defined testicular volume (TV) as the mean value of bilateral testicular volume measured by a Prader orchidometer. Serum hormone level [including follicle-stimulating hormone (FSH), luteinizing hormone (LH), total testosterone (T)] was determined before 9 a.m. Our patients were administered with human chorionic gonadotropin (hCG, Livzon Pharmaceutical Co., Zhuhai, China) due to financial reasons and/litter fertility desire. The dosage of hCG injection was 2,000 IU, twice per week. It would be adjusted according to serum T in the first 6 months. After 6 months, follow-up interval ranged from 3–6 months. Their Tanner stage, TV, serum T were recorded. Semen examination was recommended if patients were capable of masturbating. Semen examination complied to the rules of the World Health Organization laboratory manual for the examination and processing of human semen (fifth version) (17).

Conservative analysis and crystal structure prediction

The conservativeness of residue p.R240 in GNRHR protein was evaluated by sequence alignment in different vertebrates (*Homo sapiens*, *Mus musculus*, *Rattus norvegicus*, *Bos taurus*, *Ovis aries*, *Cairina moschata* and *Danio rerio*). By using a classic bioinformatic online tool, I-TASSER modeling (<https://zhanglab.ccmb.med.umich.edu/I-TASSER/>), we predicted crystal structure of GNRHR protein (both wild type and mutant type). Pattern diagram was supported by VMD software (the Beckman Institute, Illinois, USA).

Plasmids and GNRH decapeptide constructs

Plasmid (pEnter vector) of GNRHR-wild type (WT) expressing open reading frame of GNRHR (NM_000406.2) was obtained from Vigene Biosciences (Jinan, China). GNRHR-WT was fused with a C terminal Flag tag. We utilized the Quikchange II Site-Directed Mutagenesis kit (Stratagene, San Diego, CA, USA) to generate a plasmid of GNRHR-mutant type (MUT) (containing the *GNRHR* rare sequencing variant, c.719G>A, p.R240Q) based on the template plasmid. Given that function of decapeptide EHWSYGLRPG was identical to native GnRH sequence (18), we used the decapeptide obtained from

Qiangyao Biosciences (Shanghai, China) to perform functional analysis.

Cell culture and transfection

Due to deficiency of endogenous GNRHR expression, Simian virus 40 (SV40)-infected CV1 (COS-7) cells (African green monkey kidney fibroblast-like cell line, ATCC, CRL-1651) was chosen as cell model in our study (19–21). COS-7 cells were seeded at standard 6 well dishes and cultured with complete media (10% fetal bovine serum). Two microgram vector was transfected into COS-7 cells using Lipofectamine[®] 3000 Reagent (Invitrogen, Shanghai, China) at 70–80% confluency. To exclude influence of transfection, we also transfected an empty plasmid as the MOCK group.

Real-time PCR

Total RNA from transfected COS-7 cells (the GNRHR-WT, GNRHR-MUT and MOCK groups) was extracted and used for cDNA synthesis by using Rever Tra Ace[®] qPCR RT kit (Toyobo, Osaka, Japan). We performed Real-time PCR based on an MX3000P quantitative PCR system (Agilent, Santa Clara, CA, USA) by using SYBR[®] Green Realtime PCR Master Mix (Toyobo, Osaka, Japan). We performed 40 cycles of denaturing at 95 °C for 15 s, followed by annealing at 60 °C for 15 s and extension at 72 °C for 45 s. GAPDH was used as an endogenous reference. Primer sequences were as follows: GNRHR-forward: 5'-GCACGGCTGAAGACTCTAAA-3'; GNRHR-reverse: 5'-AGGCAAAGAGAAAGAAGAAGTGA-3'; GAPDH-forward: 5'-CCATCTTCCAGGAACGAGAT-3'; GAPDH-reverse: 5'-GTTGCTGACGATCTTGAGGC-3'. 2- $\Delta\Delta$ Ct analysis was conducted to calculate the mRNA level of GNRHR.

Western blots

After 24 h of transfection, protein samples were collected from the GNRHR-WT, GNRHR-MUT and MOCK groups. Cells were lysed and centrifuged at 12,000 rpm for 5 mins, and protein concentration was measured. Protein samples (25 microgram) were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (10%), and then transferred to nitrocellulose membranes. After blocking for 1 h, membranes were incubated with primary antibodies at 4 °C overnight. Herein, we used rabbit polyclonal Flag

(Signalway Antibody, 20543-1-AP, Baltimore, MD, USA) to specifically detect GNRHR fused with Flag-tag (expressed from plasmid constructs). Other primary antibodies included rabbit monoclonal ERK1/2 (Cell Signaling Technology, 137F5, Danvers, MA, USA), rabbit monoclonal p-ERK1/2^{Thr202/Tyr204} (Cell Signaling Technology, 197G2), mouse monoclonal GAPDH (Proteintech, 60004-1-Ig, Wuhan, China). Membranes were washed and then incubated with secondary antibodies, goat anti-rabbit (Boster, BA1055, Wuhan, China) or goat anti-mouse (Boster, BA1051), for 1 h at room temperature. Band Density was analyzed by Image-Pro plus software (Media Cybernetics, Rockville, MD, USA) with the corresponding GAPDH as control.

ERK1/2 activation analysis

We first examined ERK1/2 activation after GnRH decapeptide (1 μ M) stimulation (5 mins, 10 mins, 15 mins, 30 mins) in the GNRHR-WT group. The maximal response, defined as the highest level of p-ERK1/2/ERK1/2, was obtained at 5 mins. This time point was therefore chosen to perform further analysis of ERK1/2 activation. Transfected COS-7 cells were treated with GnRH decapeptide for 5 mins in the GNRHR-WT, GNRHR-MUT and MOCK groups. Protein samples were collected as previously described, and then expression of ERK1/2 and p-ERK1/2 was assessed by Western blots.

Inositol triphosphate determination

Transfected COS-7 cells were stimulated with GnRH decapeptide at concentration of 10^{-9} M, 10^{-8} M, 10^{-7} M, 10^{-6} M, 10^{-5} M for 1 h. Cell lysate was collected, and then supernatant was obtained after centrifugation at 12,000 rpm for 5 mins. We measured total inositol triphosphate (IP3) concentration by using the general IP3 ELISA kit (MyBioSource, San Diego, CA, USA) according to the standard protocol. Absorbance at 450 nm was examined with a microplate reader.

Intracellular calcium measurement

We treated transfected COS-7 cells with GnRH decapeptide (1 μ M) for 1 h as previously described (21). Medium containing GnRH decapeptide was removed, and cells were washed three times with HBSS solution (HyClone, Shanghai, China). Cells were then incubated in

4 μ M Fluo-4 AM solution (Yeasen, Shanghai, China) for 30 mins at 37 °C. Afterwards, cells were washed three times, and then incubated in HBSS solution for another 30 mins. Nuclear was stained with 6-diamidino-2-phenylindole (DAPI, Boster). Five microscopic images were randomly collected using a BX51 microscope (Olympus, Tokyo, Japan). Positive area of calcium was calculated using Image-Pro plus software.

Statistical analysis

Data analysis was performed using GraphPad Prism (GraphPad Software, San Diego, CA, USA). All the biological experiments were repeated three times independently. Chi-Square or Fisher's Exact test was used to compare the allele frequencies of variants in different cohorts. Student's *t*-tests or one-way analysis of variance was used for normally distributed data, while nonparametric tests were used for nonnormally distributed data. normally distributed data were presented as mean \pm standard deviation (SD), and nonnormally distributed data were presented as range or interquartile range. P values <0.05 were considered statistically significant.

Results

Mutation profile in IHH patients

Of the cohort of 153 IHH patients, four sporadic patients were identified to have the same heterozygous p.R240Q rare sequencing variant (RSV, MAF <1%) in *GNRHR* gene (14) (Figure 1A). The prevalence of this RSV was 2.6% (4/153), while no other *GNRHR* RSVs were found in this cohort. Additionally, these four patients also harbored additional variants of IHH causal or candidate genes. We found *PROKR2* and *CNTN2* RSVs in Patient 1, *SEMA7A* RSV in Patient 2, *TYRO3* RSV in Patient 3, as well as *AXL* and *PLXNB1* RSVs in Patient 4. All RSVs were missense and heterozygous variants. Taking into account the patient's ethnicity (East Asian), we compared variant frequency between our patient cohort and the East Asian cohort by using the gnomAD database. The results indicated that allele frequencies of the *GNRHR*, *PROKR2*, and *PLXNB1* RSVs were significantly higher in our patient cohort compared with East Asian cohort (P<0.001). However, no remarkable differences were found in frequencies of *CNTN2*, *SEMA7A*, *TYRO3*, and *AXL* RSVs (Table 1). These results suggest that the *GNRHR* RSV occurred more

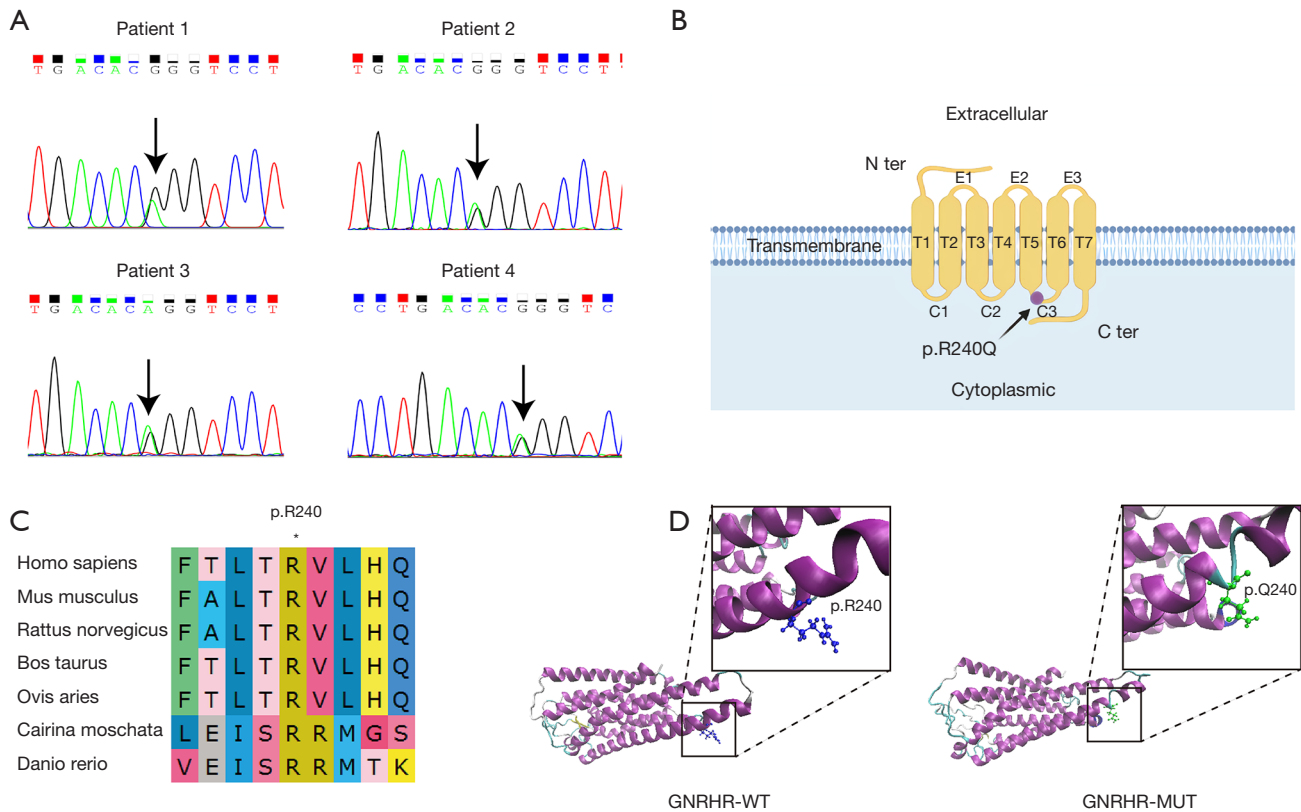


Figure 1 Sanger sequencing, prediction of structural model, conservative analysis of *GNRHR* RSV. (A) Four IHH patients harboring the *GNRHR* RSV (c.719G>A, p.R240Q) in heterozygous status. (B) Schematic representation of human *GNRHR* protein. The *GNRHR* RSV (purple dot) was located in the third cytoplasmic loop of *GNRHR*. E1-E3, T1-T7, and C1-C3 refer to the three extracellular loops, seven transmembrane domains, and three cytoplasmic loops of *GNRHR*, respectively. (C) Conservative analysis of *GNRHR* protein sequences between different seven vertebrates. The amino acid residue of p.R240 was highlighted by the asterisk. (D) Predicted structural model of *GNRHR* protein (WT and MUT). The amino acid residue of p.R240 (GNRHR-WT) and p.Q240 (GNRHR-MUT) was highlighted. Basic amino acid showed blue, and polar and neutral amino acid showed green. IHH, idiopathic hypogonadotropic hypogonadism; RSV, rare sequencing variant; WT, wild type; MUT, mutant type (containing the *GNRHR* RSV).

frequently in this Chinese IHH patient cohort than in the general population.

Clinical characteristics and outcomes

IHH is generally diagnosed during adolescence or early adulthood (1). As shown in *Table 2*, three of our four patients were diagnosed between the ages of 16 to 26 years. However, Patient 1 (43 years) was diagnosed with adult-onset IHH, since he exhibited symptoms of GnRH deficiency after sexual maturity. After detailed phenotyping, we found that Patient 2 showed anosmia but no additional phenotypes (including cryptorchidism, micropenis, cleft lip/palate, dental agenesis, optic nerve hypoplasia, sensorineural

hearing loss, etc.). With exception of Patient 2, all patients in our cohort showed testicular volume (TV) >4 mL, which suggested partial puberty development. hCG treatment significantly improved their Tanner stage, TV, serum T levels. Additionally, all patients reported the resumption of morning erections after hCG treatment. Two patients exhibited seminal spermatozoa appearance after follow-up of more than two years. No side effects (acne and gynecomastia) were observed among these patients.

Bioinformatic and expression analysis of *GNRHR* RSV

GNRHR is a member of the seven-transmembrane, G protein coupled receptor (GPCR) family in the cell

Table 1 Mutation profile in IHH patients

Patient number	Gene	Location	Zygoty	Transcript	Nucleotide change	Amino acid change	Variation frequency in patient cohort	gnomAD frequency (%)	gnomAD East Asian frequency (%)	gnomAD East Asian Homozygotes Number	P
1	<i>GNRHR</i>	chr4:68610309	Ht	NM_000406.2	c.719G>A	p.R240Q	0.013 (4/306)	0.040	0.466	0	0.000
	<i>PROKR2</i>	chr20:5283308	Ht	NM_144773.2	c.533G>C	p.W178S	0.052 (16/306)	0.018	0.260	0	0.000
	<i>CNTN2</i>	chr1:205031094	Ht	NM_005076.3	c.1075C>A	p.R359S	0.003 (1/306)	0.067	0.075	0	0.594
2	<i>GNRHR</i>	chr4:68610309	Ht	NM_000406.2	c.719G>A	p.R240Q	0.013 (4/306)	0.040	0.466	0	0.000
	<i>SEMA7A</i>	chr15:74710241	Ht	NM_003612.3	c.442C>T	p.R148W	0.003 (1/306)	0.036	0.400	0	1.000
3	<i>GNRHR</i>	chr4:68610309	Ht	NM_000406.2	c.719G>A	p.R240Q	0.013 (4/306)	0.040	0.466	0	0.000
	<i>TYRO3</i>	chr15:41865595	Ht	NM_006293.3	c.2075C>G	p.S692C	0.003 (1/306)	0.002	0.035	0	0.275
4	<i>GNRHR</i>	chr4:68610309	Ht	NM_000406.2	c.719G>A	p.R240Q	0.013 (4/306)	0.040	0.466	0	0.000
	<i>AXL</i>	chr19:41748818	Ht	NM_021913.4	c.1343G>T	p.W448L	0.003 (1/306)	0.057	0.779	4	0.570
	<i>PLXNB1</i>	chr3:48460742	Ht	NM_002673.5	c.2743T>C	p.C915R	0.003 (1/306)	0	0	0	NA

Variation frequency refers to the allele frequency of variation. P refers to the statistical significance of difference in variation frequency between the patient cohort and the East Asian cohort from the gnomAD database. IHH, idiopathic hypogonadotropic hypogonadism; Hm, Homozygous; Ht, Heterozygous; NA, not available.

membrane, and the *GNRHR* RSV is located in the third cytoplasmic loop of *GNRHR* (Figure 1B). Conservative analysis revealed that residue of p.R240 was highly conserved across seven different vertebrates (Figure 1C). In crystal structure prediction, replacement of arginine (strongly basic) with glutamine (neutral) changed the polarity of residue. The structure of this site has changed from alpha helix to curl. Glutamine has four fewer hydrogen atoms and two fewer nitrogen atoms (Figure 1D), which means that the replacement likely altered interactions with amino acids at other sites, and may further impair their function. Expression analysis showed that mRNA level of *GNRHR* in the *GNRHR*-WT group was mildly higher than that in *GNRHR*-MUT group (containing the *GNRHR* RSV), but the difference was not significant. Similarly, no significant difference was observed in protein comparison (Figure 2A,B,C).

GNRHR RSV affected ERK1/2 activation

One of the major signal transduction pathways of *GNRHR* is the MAPK cascade (9,22,23). To figure out the effect of *GNRHR* RSV on ERK1/2 activation, we first set different time points to find the optimal time of activation in the *GNRHR*-WT group. As shown in Figure 2D,E, the level

of p-ERK1/2/ERK1/2 was highest at 5 min relative to 10 min, 15 min and 30 min. The level of p-ERK1/2/ERK1/2 gradually decreased as time elapsed. We subsequently used 5 mins for the duration of stimulation. Compared with the *GNRHR*-WT group, the level of p-ERK1/2/ERK1/2 was markedly lower in the *GNRHR*-MUT group ($P < 0.05$, Figure 2E,G). The data thereby indicate that the *GNRHR* RSV impairs ERK1/2 activation to some extent.

GNRHR RSV affected IP3/calcium pathway

Because of IP3-mediated calcium mobilization in downstream of *GNRHR* (9,21), we examined IP3 accumulation and calcium mobilization to analyze the downstream effects of *GNRHR* RSV. As for IP3 accumulation, there was a remarkable difference between the *GNRHR*-WT and *GNRHR*-MUT groups ($P < 0.001$), with the concentration of the GnRH decapeptide ranging from 10^{-8} M to 10^{-5} M (Figure 3A). For calcium mobilization, the highest calcium level was shown in the *GNRHR*-WT group. Calcium level partially, but significantly decreased in the *GNRHR*-MUT group than the *GNRHR*-WT group ($P < 0.05$) (Figure 3B,C). Therefore, our result suggests that the *GNRHR* RSV partly damages IP3/calcium signal.

Table 2 Clinical characteristics of IHH patients harboring the *GNRHR* RSV.

	Patient 1	Patient 2	Patient 3	Patient 4
At baseline				
Age at diagnose (year)	43	26	21	16
Sex	Male	Male	Male	Male
BMI (kg/m ²)	24.677	24.419	22.039	21.718
Diagnose	AOIHH	KS	nIHH	nIHH
Olfactory disorder	–	Anosmia	–	–
Cryptorchidism or cryptorchidism history	–	–	–	–
Previous testosterone/gonadotropin treatment	–	–	–	–
Pubic hair Tanner stage	5	1	1	1
Genital Tanner stage	5	2	3	3
Testicular volume [†] (mL)	12	2	4.5	12
Basal FSH (mIU/mL)	0.81	2.27	2.61	0.19
Basal LH (mIU/mL)	1.06	0.25	0.54	0.00
Basal T (mIU/mL)	0.41	0.54	0.15	0.01
After treatment				
Follow-up time (month)	25	41	12	2
Drug injection dose (U/per week)	hCG 4000	hCG 4000	hCG 4000	hCG 4000
Morning erection	+	+	+	+
IIEF-5 score	21	NA	NA	NA
Pubic hair Tanner stage	5	5	4	1
Genital Tanner stage	5	4	3	3
Testicular volume (mL)	20	12	10	12
Therapeutic T (ng/mL)	2.71	4.25	3.16	1.82
Sperm concentration (10 ⁶ /mL)	13.56	4.01	NA	NA
Sperm forward progression (%)	46.55	1.34	NA	NA

[†]Testicular volume refers to the mean value of bilateral testicular volume. IHH, idiopathic hypogonadotropic hypogonadism; AOIHH, adult-onset idiopathic hypogonadotropic hypogonadism; KS, Kallmann syndrome; nIHH, normosmic idiopathic hypogonadotropic hypogonadism; RSV, rare sequencing variant; BMI, body mass index; FSH, follicle-stimulating hormone; LH, luteinizing hormone; T, testosterone; IIEF, International index of erectile function; NA, not applicable.

Discussion

In this study, we described a RSV in *GNRHR* gene and associated clinical manifestation. We identified four sporadic patients carrying the same heterozygous *GNRHR* RSV (p.R240Q). The prevalence of this *GNRHR* RSV was 2.6%. Its allele frequency was found to be exceptionally higher than that in general population. Functional analysis has demonstrated that this RSV caused loss of function

(LOF) to *GNRHR* protein. It is of value to investigate the effect of the RSV, although the heterozygous state may be a non-functional factor in human, it homozygous/compound state could possibly have a pathogenic effect.

Heterozygous changes in *GNRHR* gene were enriched in IHH patients compared with controls. A large genetic screening for 863 probands with different forms of GnRH deficiency found that, 24 patients with monoallelic

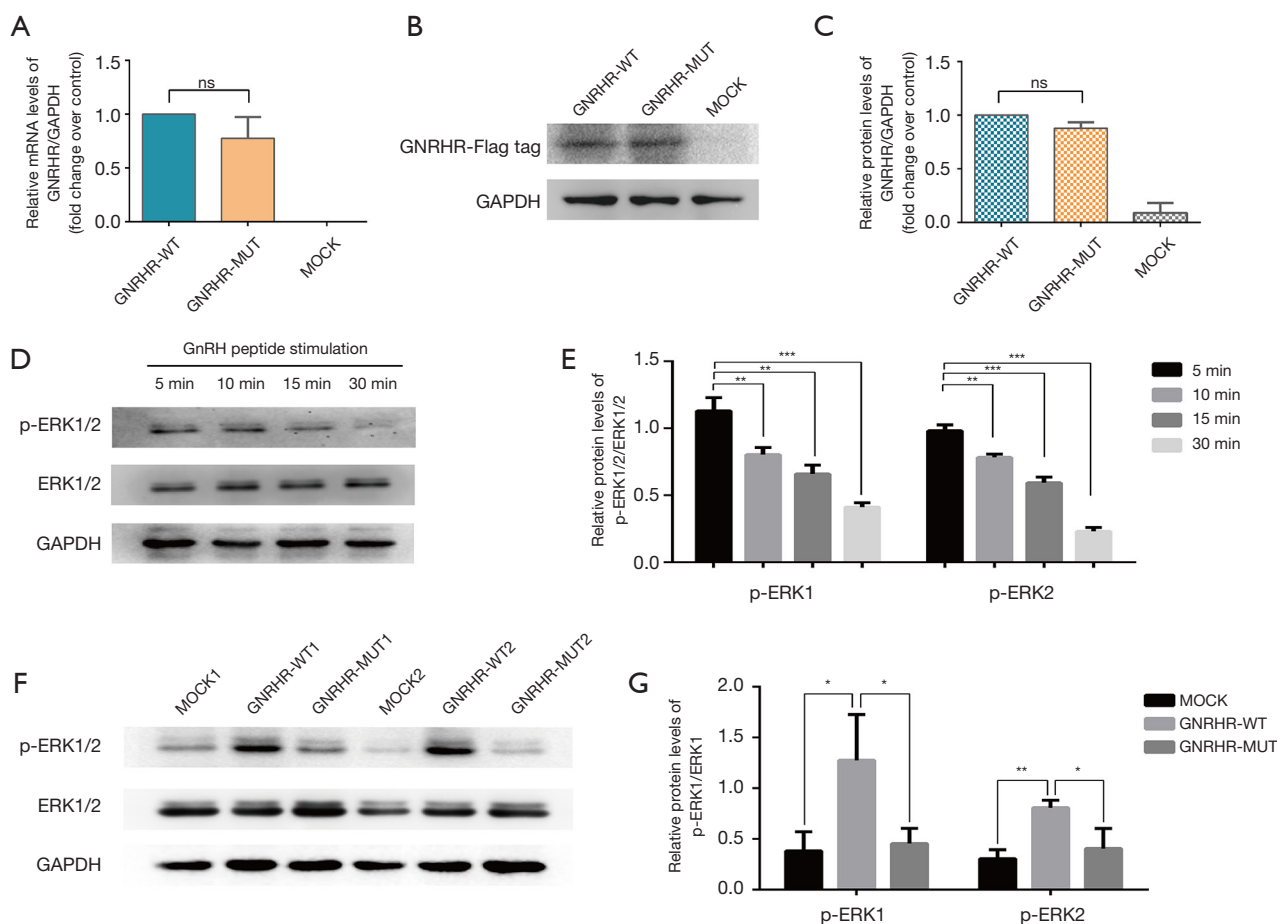


Figure 2 Analysis of GNRHR expression and ERK1/2 activation under stimulation of GnRH decapeptide. (A) Quantitative analysis of GNRHR mRNA level in the GNRHR-WT, GNRHR-MUT, and MOCK groups. (B) Representative western blot results of GNRHR expression in the GNRHR-WT, GNRHR-MUT, and MOCK groups. (C) GNRHR expression with GAPDH as the loading control in the GNRHR-WT, GNRHR-MUT, and MOCK groups. (D) Representative western blot results of p-ERK1/2 and ERK1/2 in the GNRHR-WT group under the stimulation of GnRH decapeptide at different time points. (E) Relative protein level of p-ERK1/ERK1 and p-ERK2/ERK2 in the GNRHR-WT group at different time points. (F) Representative western blot results of p-ERK1/2 and ERK1/2 in the GNRHR-WT, GNRHR-MUT, and MOCK groups under stimulation for 5 min. (G) Relative protein level of p-ERK1/ERK1 and p-ERK2/ERK2 in the GNRHR-WT, GNRHR-MUT, and MOCK groups. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$. RSV, rare sequencing variant; WT, wild type; MUT, mutant type (containing the *GNRHR* RSV); ns, not significant. The data were obtained from three biological replicates.

GNRHR mutations featured a broad phenotypic spectrum, ranging from Kallmann syndrome/nIHH, to attenuated GnRH deficiency (AOIHH, hypothalamic amenorrhea, or constitutional delay of puberty) (12). Intriguingly, researchers revealed that the heterozygous p.Q106R and R262Q RSVs were verified to be partial loss-of-function, and the heterozygous R139H RSV was complete loss-of-function (6,12). A recent study showed that two KS patients and one nIHH patient carried this heterozygous p.Q106R RSV, and they did not carry any other 36 IHH-

associated gene variants (11). Therefore, these evidence indicates that heterozygous changes in *GNRHR* are not just a non-functional factor, but may lead to reproductive pathogenicity (11). Given the autosomal recessive inheritance, our study carefully suggests that the p.R240Q RSV found in four IHH patients may play a limited role in the process of morbidity. One possibility is that other mutations with unknown function on these patients caused the disease to occur. An additional possibility is that decreased gene dosage caused by monoallelic *GNRHR*

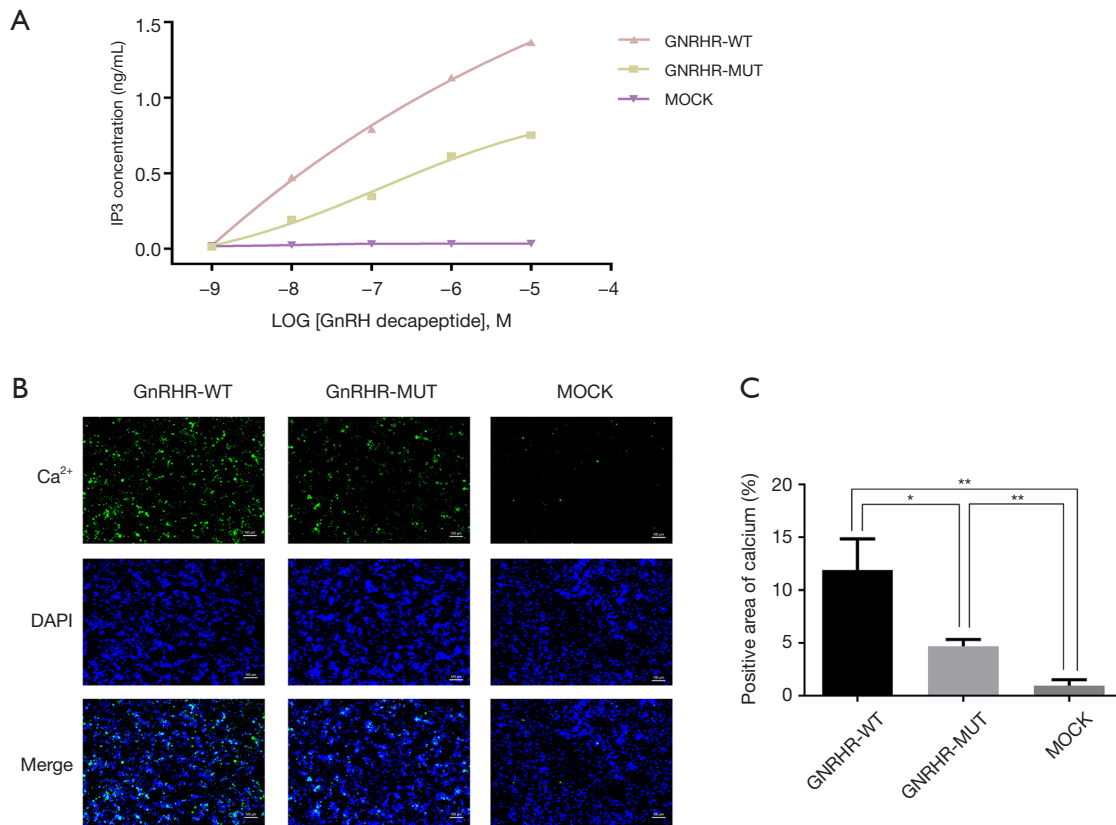


Figure 3 *GNRHR* RSV negatively affected intracellular IP3 and calcium level. (A) The intracellular IP3 concentration under stimulation of different concentrations of GnRH decapeptide [LOG (GnRH decapeptide), M = -9, -8, -7, -6, -5] in the GNRHR-WT, GNRHR-MUT, and MOCK groups. (B) Representative results of intracellular calcium level in the GNRHR-WT, GNRHR-MUT, and MOCK groups (stained with Fluo-4 AM and DAPI) (magnification $\times 100$, Scale bars = 100 μm). (C) Semiquantitative analysis of positive area of intracellular calcium in the GNRHR-WT, GNRHR-MUT, and MOCK groups. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$. RSV, rare sequencing variant; WT, wild type; MUT, mutant type (containing the *GNRHR* RSV); IP3, Inositol trisphosphate. The data were obtained from three biological replicates.

mutation, makes individuals susceptible to environmental, behavioral, or psychosocial burdens on GnRH secretion (11,24).

Although *GNRHR* mutations were reported in normosmic IHH patients (25), we have demonstrated that the four patients showed phenotypic diversity. Patient 1, 3 and 4 had normal sense of smell, whereas Patient 2 were diagnosed with KS (containing olfactory disorder). Patient 1 displayed GnRH deficiency in adulthood, while the other three patients displayed that in pubertal development. At baseline, there were different levels of testicular development and hormone profiles. Testicular volume of Patient 3 and 4 were both above >4 mL, suggesting partial spontaneous puberty development (26), whereas TV of

Patient 2 was 2 mL. Moreover, serum FSH levels of Patient 2 and 3 were at the lower level of normal range, but the rest two patients did not showed that. Consistently, other studies also proved this phenotypic diversity in patients with *GNRHR* mutations, ranging from complete IHH to constitutional delay of puberty (7,12,21).

Digenic and oligogenic inheritance are commonly observed in IHH (14,27). Herein, four patients with the *GNRHR* RSV harbored additional genetic abnormalities, including known IHH causal genes *PROKR2*, *SEMA7A*, *AXL* RSVs, and candidate genes *CNTN2*, *TYRO3*, and *PLXNB1* RSVs. Except for the defined pathogenicity of *PROKR2* RSV (p.W178S) (28,29), the role of remaining RSVs are currently not well understood. Of note, all

four patients in our cohort showed good responses to hCG treatment. After follow-up of over 2 years, fertility outcomes of two patients were improved, with sperm concentration of 13.56 and 4.01×10^6 /mL, respectively. Early studies indicated that TV >4 mL was a positive predictive factor for sperm production after receiving gonadotropin treatment (1,30,31). Given that the rest two patients also had TV >4 mL, it is plausible that hCG treatment may induce spermatogenesis in further follow-up.

It is well established that GnRH receptor is a member of the G protein coupled receptors (GPCRs), and highly expressed in the anterior pituitary gland, breast, ovary, and prostate (32). Mutations in GPCRs disrupt a variety of physiological processes, including protein biosynthesis, folding and transport, ligand binding, receptor activation, and interaction with G protein (10,33). Seventy-five percent of *GNRHR* variants reside in transmembrane domains (10). Our genetic analysis found that the RSV (p.R240Q) was located in the third cytoplasmic loop of GNRHR. As described in earlier studies (6,10,34-36), there are two more *GNRHR* variants (p.R262Q and p.L266R) reported in this loop. These variants both disrupted normal GNRHR expression and downstream signals. However, in our study, the p.R240Q RSV had no effect on expression, but negatively affected downstream signals, strongly suggesting that pathogenicity of this RSV may be milder than that of p.R262Q and p.L266R for the GNRHR protein.

According to severity of impaired function, we divided *GNRHR* variants into two subtypes. Complete loss-of-function (cLOF) is defined as no measurable in vitro signaling, while partial loss-of-function (pLOF) is defined when maximal signaling intensity is at least 10% of wild-type signaling (10). Protein analysis showed that the level of ERK1/2 phosphorylation in the GNRHR-MUT group was about 36–50% of that in the GNRHR-WT group. IP3 accumulation and calcium mobilization of GNRHR-MUT was approximately 20–54% of GNRHR-WT. Therefore, the *GNRHR* RSV is classified as a pLOF mutation. Moreover, our finding suggests that pLOF mutation may be associated with mild phenotypic forms, such as partial testicular development and normal FSH levels.

There are several limitations in this study. First, due to the geographic and economic reasons, it is difficult to obtain blood samples from patients' family and ascertain their genotypes. Second, each patient harbored other gene

mutations, we could not exclude the role of other possible pathogenic variants in the pathogenesis of IHH. Third, the intervals of follow-up review ranged from 3 to 6 months. A shorter and more consistent follow-up would be better for clinical management. Additionally, knowledge of IHH genes is rapidly progressing, our panel sequencing needs to be updated and more new causal genes should to be added into the panel.

Conclusions

In conclusion, the present study supports the loss of function of the *GNRHR* RSV (p.R240Q), and further expands its mutation and phenotypic spectrum. Monoallelic defects in *GNRHR* was found in Chinese IHH cohort, and it should be paid attention in this research field.

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Footnote

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Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study protocol (IRB ID: 20150302) (including collection of blood samples) was approved by the Ethics Committee of Huazhong

University of Science and Technology in compliance with the Declaration of Helsinki (as revised in 2013). Informed consent was signed with each patient.

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