

## NOBOX Homeobox Mutation Causes Premature Ovarian Failure

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*NOBOX* (newborn ovary homeobox gene) is an oocyte-specific homeobox gene that plays a critical role in early folliculogenesis and represents a candidate gene for nonsyndromic ovarian failure. We investigated whether mutations in the *NOBOX* gene cause premature ovarian failure (POF). We sequenced the *NOBOX* gene in 96 white women with POF and discovered seven known single-nucleotide polymorphisms and four novel variations, two of which, p.Arg355His and p.Arg360Gln, cause missense mutations in the homeobox domain. Electrophoretic mobility shift assay (EMSA) confirmed that the missense mutation, p.Arg355His, disrupted *NOBOX* homeodomain binding to *NOBOX* DNA-binding element (NBE) and had a dominant negative effect on the binding of wild-type *NOBOX* to DNA. Our findings demonstrate that *NOBOX* mutations can cause POF.

In premature ovarian failure (POF [MIM 311360]), ovaries cease to mature oocytes before 40 years of age. The condition is characterized by secondary amenorrhea, infertility, hypoestrogenism, and elevated gonadotropin serum levels (FSH > 40 IU/liter).<sup>1</sup> POF is heritable in up to 30% of patients and is genetically heterogeneous.<sup>2,3</sup> Mechanisms long invoked in the pathogenesis of POF include abnormalities of the X chromosome or autosomes and autoimmune, infectious, and environmental causes.<sup>4,5</sup> For some cases, a genetic basis has been shown, and, for genes such as *FMR1* (MIM 309550),<sup>6,7</sup> *FSHR* (MIM 136435),<sup>8-10</sup> *POF1B* (MIM 300603),<sup>11</sup> *FOXL2* (MIM 605597),<sup>12,13</sup> and *BMP15* (MIM 300247),<sup>14</sup> functional data support causation. Because human ovaries are not easily accessible, our knowledge of ovarian development is largely derived from animal models.<sup>15</sup> Mouse knockouts and naturally occurring mutations in sheep have been useful in identifying candidate genes for ovarian failure.<sup>16-19</sup>

Human and mouse *NOBOX* genes (newborn ovary homeobox [MIM 610934]) are preferentially expressed in oocytes and encode a homeobox transcriptional regulator.<sup>20,21</sup> *Nobox* plays a crucial role during early folliculogenesis in the mouse.<sup>22</sup> Disruption of the mouse *Nobox* gene causes nonsyndromic ovarian failure in females, whereas males are unaffected.<sup>23</sup> *NOBOX* expression in adult human tissues mimics that in mice, with preferential expression in the human gonads.<sup>21</sup> Expression within the human ovary is oocyte specific, as observed from primordial follicle through metaphase II (MII) oocytes. Mouse *NOBOX* homeodomain binds TAATTG,<sup>24</sup> and we have shown elsewhere that *NOBOX* binds such elements in *Gdf9* (MIM 601918) and *Pou5f1* (MIM 164177) promoters.<sup>25</sup> In the current study, we report two novel missense mutations in the *NOBOX* homeodomain in women

with POF and show that p.Arg355His mutation can disrupt *NOBOX* homeodomain binding to DNA.

Our study subjects comprised 96 white women from the United States who had POF and had been collected at Baylor College of Medicine since 2001. Recruitment criteria comprised cessation of menstrual cycles before 40 years of age and at least two serum FSH concentrations >40 IU/liter. Women with chromosomal abnormalities were excluded. Two hundred and seventy-eight white women who denied having any medical problems were used as controls. Informed consent for molecular studies was obtained from all subjects. The study was approved by the Institutional Review Board of Baylor College of Medicine.

Peripheral blood was obtained and genomic DNA extracted. *NOBOX*-specific primers were designed according to the human *NOBOX* sequence (GenBank accession number NM\_001080413 and Ensembl accession number ENST00000389325) and are presented in table 1.

All 10 exons and exon-intron boundaries of the *NOBOX* gene were amplified using PCR. PCR conditions are available on request. PCR products were sequenced directly on an automated sequencer, ABI Prism Sequencer 3730XL (Applied Biosystems).

Our study revealed 11 sequence variants in the coding region of *NOBOX*: 7 known SNPs and 4 novel variations (fig. 1A). All sequence variants were confirmed by three independent PCR and sequencing reactions, followed by sequencing in forward and reverse directions. Details are provided in table 2.

The seven known SNPs included three intronic variants (*rs757388*, *rs11769847*, and *rs11979528*), two synonymous variants (c.42T→C and c.262C→T), and two non-synonymous variants (p.Gly482 Ser and p.Phe228Leu). Of

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**Table 1. PCR Primers Used for Amplification of Human NOBOX Gene**

Primer	Sequence (5'→3')	Reference Sequence <sup>a</sup>
NOBOX 1F	ATTTAAAGACAAGCTCGAGTATC	NM_001080413
NOBOX 1R	CAAGAGTCTCAGTGTATGGG	
NOBOX 2F	TATCTGACCAGCCCTCCGACTTT	NM_001080413
NOBOX 2R	ACTGCTGGAATTACAGGCGTGAG	
NOBOX 3F	ATGCTCTTGCTCGGCTGCTGGA	NM_001080413
NOBOX 3R	AGCTTGACTATTGTGAGGATT	
NOBOX 4F	TGCTAAGGTCAGTGGATGTTGGC	NM_001080413
NOBOX 4R	CCGGGACGAAGTGACATAC	
NOBOX 5F	TGTAGCACCCAAGAGCGAGAA	NM_001080413
NOBOX 5R	TGCAGTGCTCTCTCTCTAATG	
NOBOX 6-7F	GCAACAGCCAGGACCTAAGC	NM_001080413
NOBOX 6-7R	GTCACCTCCACCTCCATCAAACA	
NOBOX 8F	TCTACCATTACAGCGATGCCAA	NM_001080413
NOBOX 8R	TGTGCTCTCTCAGTAAACC	
NOBOX 9F	GGGACTCCGCTACTGTGGT	ENST00000389325
NOBOX 9R	CGAGGGAGAAGAGCTTAATAG	
NOBOX 10F	GTCCTAAGCTGCGTCTATGTG	ENST00000389325
NOBOX 10R	CGAGCCCAATCTATCCCA	

<sup>a</sup> NM\_001080413, Genbank. ENST00000389325, Ensembl.

the four novel variations, one was a synonymous variant (c.66T→C) and three were nonsynonymous exonic variants: c.1354G→A (p.Asp452Asn), c.1064G→A (p.Arg355His), and c.1079G→A (p.Arg360Gln). p.Asp452Asn was also found in five controls at a frequency that did not differ significantly from that in the POF sample (Fisher's exact test  $P > .05$ ). p.Arg355His and p.Arg360Gln were located in the conserved homeodomain region (fig. 1A–1C). p.Arg360Gln was also present in two controls at a frequency that was not statistically different from that in women with POF (Fisher's exact test  $P > .05$ ). By contrast, p.Arg355His was not present in the controls. p.Arg355His mutation was found in a 35-year-old woman whose FSH level was 103 IU/liter. Her menarche was at age 11 years, and she entered menopause at age 32 years. She was the

mother of two healthy children and lacked overt somatic anomalies. She was the only child of a woman who conceived her at 26 years of age and who entered menopause at 42 years of age.

The protein sequence of human NOBOX is 92% identical to the corresponding homeodomain region in the mouse.<sup>20</sup> Both mutations, p.Arg355His and p.Arg360Gln, were located within the homeodomain region that is perfectly conserved among species ranging from zebrafish to humans. Since a homeodomain is capable of mediating critical protein-DNA and protein-protein interactions, and since p.Arg355His was not present in control women, we hypothesized that this novel mutation disrupted NOBOX homeodomain binding to DNA.

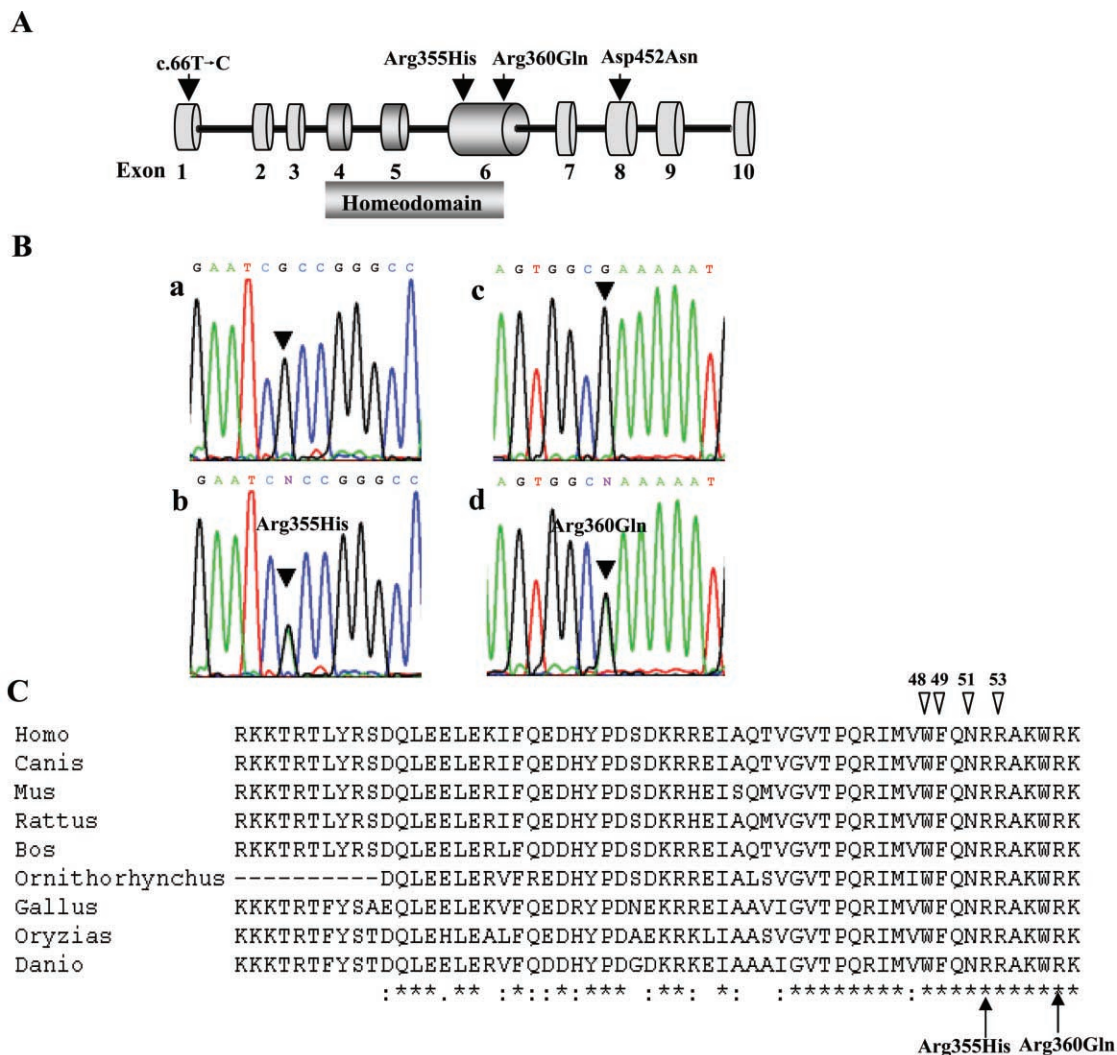
We engineered p.Arg355His and p.Arg360Gln missense mutations into the mouse homeodomain to test whether these mutations will disrupt mouse NOBOX homeodomain binding to the NOBOX DNA-binding element (NBE) we identified elsewhere.<sup>25</sup> The mouse homeobox was subcloned into the pET41b expression vector to generate a fusion protein with glutathione-S-transferase (GST-NXHD). Mouse missense mutations corresponding to human p.Arg355His and p.Arg360Gln (mouse p.Arg186His and p.Arg191Gln) were generated using the QuikChange multi site-directed mutagenesis kit (Stratagene) with oligonucleotides GTGTGGTTTCAGAACCACAGGGCAAAGTGGAGA and CGCAGGGCAAAGTGGCAGAAAGTGGAGAACTG. We transformed BL21-pLysS *Escherichia coli* (Stratagene) with GST-NXHD (wild type), GST-R186H (containing mutation p.Arg186His), and GST-R191Q (containing mutation p.Arg191Gln) constructs, induced protein expression, and purified the GST-NXHD, GST-R186H, and GST-R191Q fusion proteins on GST-bind resin (Novagen). The proteins were dialyzed three times in PBS to remove excess glutathione and were quantified with BSA by use of the Lowry assay.

**Table 2. NOBOX Sequencing Results in 96 White Women with POF**

Sequence Variation	Location	Amino Acid Variation	dbSNP ID	Allele Frequency (%)					
				Patients with POF			Controls		
				Wild Type	Heterozygote	Homozygote	Wild Type	Heterozygote	Reference <sup>a</sup>
c.42T→C	Exon 1	Synonymous	rs1208179	71.9	28.1	0			d, e
c.66T→C	Exon 1	Synonymous	Novel	98.96	1.04	0	100	0	d, e
c.262C→T	Exon 3	Synonymous	rs727714	11.5	45.8	42.7			d, e
c.1064G→A	Exon 6	p.Arg355His	Novel	99.0	1.01	0	100	0	d, e
c.1079G→A <sup>b</sup>	Exon 6	p.Arg360Gln	Novel	99.0	1.01	0	99.3	.7	d, e
c.1154+11T→C	Intron 6		rs757388	11.5	45.8	42.7			d, e
c.1155-22G→A	Intron 6		rs11769847	11.5	45.8	42.7			d, e
c.1354G→A <sup>b</sup>	Exon 8	p.Asp452Asn	Novel	99.0	1.01	0	94.8	5.2	d, e
c.1444G→A	Exon 8	p.Gly482Ser	rs2525702	72.9	26.0	1.1			d, e
c.603-51G→T	Intron 8		rs11979528	29.2	44.8	26.0			f, g
c.682T→C	Exon 9	p.Phe228Leu	rs2699503	14.6	47.9	37.5			f, g

<sup>a</sup> d = GenBank accession number NM\_001080413. e = NCBI protein database accession number XP\_001134420. f = Ensembl accession number ENST00000389325. g = Ensembl accession number ENSP00000373976.

<sup>b</sup> Allelic frequencies between the general population and patients with POF show no significant differences (Fisher's exact test  $P > .05$ )

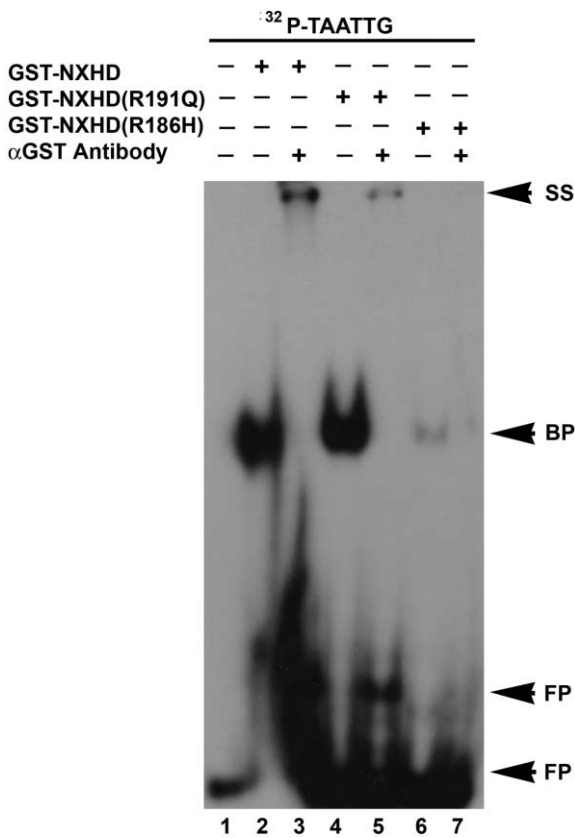


**Figure 1.** Identification of novel variants in the *NOBOX* gene. *A*, Schematic diagram of the *NOBOX* gene, showing three novel non-synonymous variations (exon 6 and exon 8) and one synonymous variant (exon 1) found in the present study. Exons are a composite of human sequence NM\_001080413 (GenBank) and ENST00000389325 (Ensembl). The homeodomain region is coded by exons 4–6. *B*, Electropherogram showing wild-type (*a* and *c*) and mutant (c.1064G→A [Arg355His] and c.1079G→A [Arg360Gln]) sequences. Arrows indicate the G→A change in POF subject 23 (*b*, Arg355His) and POF subject 34 (*d*, Arg360Gln). *C*, *NOBOX* homeodomain alignment among different species. Arrow heads indicate highly conserved amino acids highly conserved among homeodomains. The two missense mutations (p.Arg355His and p.Arg360Gln) found in this study are indicated by arrows. Asterisks (\*) indicate amino acid residues that are conserved in all the species.

We performed electrophoretic mobility shift assay (EMSA) as described elsewhere.<sup>25</sup> In brief, NBE (containing *NOBOX*-binding consensus sequence TAATTG) was labeled by end-filling annealed primers with [ $\alpha$ -<sup>32</sup>P] dCTP and Klenow polymerase (Invitrogen) at room temperature. The top and bottom strand oligonucleotides were 5'-ACG-AGCTACCTACTTAATTGACGT-3' and 5'-ACAGTACG-CGTTCAACGTC-3', respectively. Binding reactions were conducted by incubating <sup>32</sup>P-labeled probe (250,000 cpm/reaction) individually with 50 ng of purified GST-NXHHD, GST-R186H, and GST-R191Q. Polyclonal anti-GST antibodies (Amersham Biosciences) were used to supershift

DNA-protein complexes. Binding reactions were resolved on a 4% polyacrylamide gel. The gel was then fixed, dried, and exposed to Kodak BioMax XAR Film.

As shown in figure 2 (lanes 2 and 4), GST-NXHHD or GST-R191Q protein bound radiolabeled NBE with similar affinity. In contrast, GST-R186H protein showed a dramatic decrease in the NBE-binding capacity (fig. 2, lane 6). In the supershift assay, addition of anti-GST antibodies decreased the mobility of the GST-NXHHD-NBE and GST-R191Q-NBE complex and, as expected, diminished the total amount of the DNA-protein complex (fig. 2, lanes 3, 5, and 7). These results show that R186H mutation dis-



**Figure 2.** GST-R186H binds to the TAATTG sequences with low affinity. <sup>32</sup>P-labeled TAATTG was incubated with purified recombinant GST-NXHD, GST-R191Q, or GST-R186H. <sup>32</sup>P-labeled TAATTG formed a strong DNA-protein complex with the recombinant GST-NXHD (lane 2) and GST-R191Q protein (lane 4), but not with GST-R186H (lane 6). The DNA-protein complex was supershifted when incubated with polyclonal antibodies against GST (lanes 3, 5 and 7). The presence or absence of recombinant protein and/or antibody is indicated above each lane with a plus (+) or minus (-) sign, respectively. Arrows indicate DNA-protein complex (BP), supershifted complex (SS) and free probe (FP). Oligonucleotide sequence containing TAATTG is 5'-ACG AGC TAC CTT ACT TAA TTG GAC GTT GAA CGC GTA CTG T-3'.

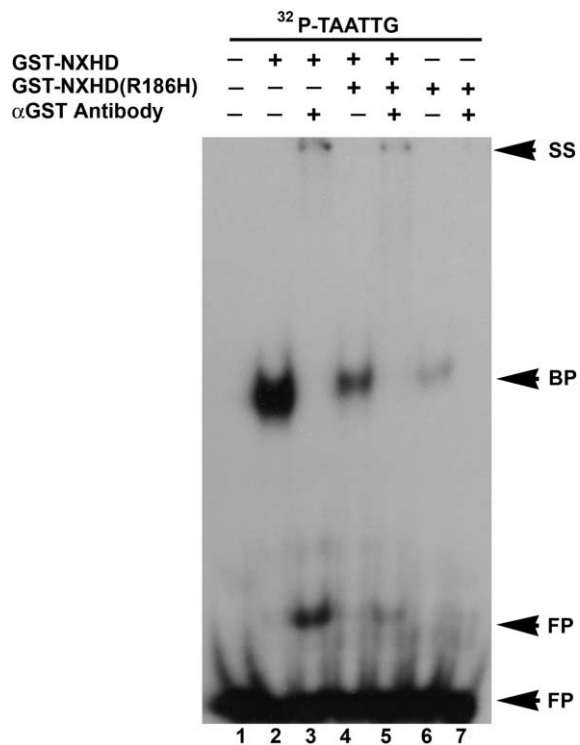
rupted NOBOX homeodomain binding, whereas R191Q did not exhibit a similar effect.

We also examined whether the R186H mutation interfered with the DNA-binding capacity of the wild-type NOBOX homeodomain protein. Equivalent amounts of GST-NXHD and GST-R186H (50 ng) were coincubated (fig. 3), and the mixture of wild-type and R186H homeodomains exhibited weaker binding to NBE than did wild type alone (fig. 3, lanes 4 and 2). These experiments indicate that the R186H mutation has a dominant negative effect on the wild-type NOBOX homeodomain and that it causes diminished binding affinity for the NBE. This effect was not observed with the R191Q missense mutation.

In this study of 96 white women with POE, we found two missense mutations in the NOBOX homeodomain,

one of which, p.Arg355His, is absent in the control population and disrupts NOBOX homeodomain binding to NBE in a dominant negative fashion. Mice that are null for *Nobox* are infertile and lose oocytes rapidly during ovarian development. *Nobox* is therefore a critical oocyte prosurvival factor. Partial deficiency in NOBOX function in humans can conceivably lead to a more gradual loss of oocytes in women and can cause POF after the onset of puberty.

Homeodomains are generally composed of 60 aa, as is predicted for NOBOX. The p.Arg355His missense mutation is located within the highly conserved portion of the NOBOX homeodomain at position 52 (Arg52) (fig. 1C). Several amino acids are conserved in the majority of homeodomains. These include Trp48 (W), Phe49 (F), Asn51 (N), and Arg53 (R) within the homeodomain (fig. 1C); all reside in helix III, which has critical implications for DNA binding and overall stability of the tertiary structure of the homeodomains.<sup>24</sup> Prior studies have demonstrated that substitution of glutamine for the conserved aspara-



**Figure 3.** Dominant negative effect of GST-R186H. Radiolabeled NBE (<sup>32</sup>P-TAATTG) was incubated individually with purified recombinant GST-NXHD, GST-R186H, or a 50/50 mixture of both homeodomains. GST-R186H reduced GST-NXHD binding affinity to the <sup>32</sup>P-labeled TAATTG (lane 4). The specificity of the DNA-protein complexes was confirmed by polyclonal antibodies against GST. The presence or absence of recombinant protein and/or antibody is indicated above each lane with a plus (+) or minus (-) sign, respectively. Arrows indicate DNA-protein complex (BP), supershifted complex (SS) and free probe (FP).

gine residue at position 51 abolished NOBOX homeodomain binding to NBES.<sup>25,26</sup>

Despite histidine being a polar amino acid like arginine, p.Arg355His mutation, when engineered into a mouse homeodomain, disrupts NOBOX binding to DNA. Arg355 in the NOBOX homeodomain is conserved from zebrafish to humans and is a critical residue for NOBOX homeodomain binding to the DNA. Homeodomains are also known to be critical in protein-protein interactions, involving either homeodomains or other protein domains.<sup>27</sup> Direct protein-protein interactions are necessary for transcriptional activity.<sup>28,29</sup> The dominant negative effect also suggests that the NOBOX homeodomain may function as a dimer. The p.Arg360Gln missense mutation, when engineered into a mouse homeodomain, does not disrupt protein-DNA interaction in the EMSA assay, was also found in the control population, and is consistent with being a novel SNP.

Our study shows that mutations within *NOBOX* can cause human POF. The only previous study involved 35 Japanese women with POF and did not find mutations in the *NOBOX* homeodomain.<sup>30</sup> By contrast, our results indicate that mutations within *NOBOX* likely account for POF in a small subset of women. There are numerous oocyte-specific genes, each of which may be responsible for some cases of POF.<sup>31</sup> Future high-throughput sequencing of such genes in women with idiopathic ovarian failure should provide a better idea of the contribution that oocyte-specific genes make to nonsyndromic POF.

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## Web Resources

Accession numbers and URLs for data presented herein are as follows:

Ensembl, <http://www.ensembl.org/> (for *NOBOX* [accession number ENST00000389325])

dbSNP, <http://www.ncbi.nlm.nih.gov/SNP/> (for IDs shown in table 2)

GenBank, <http://www.ncbi.nlm.nih.gov/Genbank/> (for *NOBOX* [accession number NM\_001080413])

NCBI Protein Database, <http://www.ncbi.nlm.nih.gov/sites/entrez?db=protein&cmd=search&term=> (accession number XP\_001134420)

Online Mendelian Inheritance in Man (OMIM), <http://www.ncbi.nlm.nih.gov/Omim> (for POF, *NOBOX*, FMR1, POF1B, FSHR, FOXL2, BMP15, GDF9, and POU5F1)

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