

**Postdoc position**  
**in Human Reproduction Genetics**  
**in Heidelberg**  
**(Fulltime for 3 years; DFG grant):**

**Major Research topic:**

**“Molecular control signature of *FMR1*/FMRP  
expression in human granulosa cells functional  
during folliculogenesis and for ovarian reserve”**

Research projects in my research group (AG “Genetics of Human Folliculogenesis”) at *Division of Reproductive Genetics (head: Prof. Dr. P.H. Vogt)* are concentrated on the molecular analysis of genetic and epigenetic regulatory mechanisms controlling *FMR1*/FMRP expression during human folliculogenesis and its putative distortion in some well-known female germ line disorders such as Premature Ovarian Insufficiency/Failure (POI/POF) syndrome and Poor Ovarian Response (POR).

Our aim is to identify the molecular tools functional in genetic germ line networks as the mTOR/AKT/FOXO signal pathway associated with the regulation of *FMR1*/FMRP expression in the female germline cells (oocytes, granulosa cells, theca cells). Our results will help not only to understand much better how these genes/proteins impact **human female fertility**, but also to develop novel starting points for improved molecular diagnostic and therapeutic tools for **infertility treatment and assisted reproductive techniques (ART)**.

[Related Publications:](#)

- Genetische Ursachen der prämaternen Ovarialinsuffizienz  
**Rehnitz J**, Strowitzki T, Vogt PH. Gynäkologische Endokrinologie. Ausgabe 04/2018
- The *FMR1* gene, infertility, and reproductive decision-making: a review.  
Pastore LM, Johnson J.; Front Genet. 2014 Jul 7;5: 195. Review
- Variable expression of the *Fragile X Mental Retardation 1 (FMR1)* gene in patients with premature ovarian failure syndrome is not dependent on number of (CGG)<sub>n</sub> triplets in exon 1. **Schuetzler J**, Peng Z, Zimmer J, Sinn P, von Hagens C, Strowitzki T, Vogt PH: Hum Reprod. 2011 May;26(5):1241-1251.
- *FMR1* and *AKT/mTOR* signaling pathways: potential functional interactions controlling folliculogenesis in human granulosa cells. **Rehnitz J**, Alcoba DD, Brum IS, Hinderhofer K, Youness B, Strowitzki T, Vogt PH. Reprod Biomed Online. 2017 Nov;35(5):485-493.

## Your profile

- PhD degree in molecular biology, biochemistry or equivalent (preferable but not essential with topic centered in *Human Reproduction Biology*)
- Splendid expertise in all molecular biology techniques and extensive experience in human cell culture work. Especially: Western blotting, Immunoprecipitations, ChIP, gene expression analysis, PCR, qPCR, cloning, sequencing, ELISA, Fluorescent microscopy, Next-Generation-Sequencing, CRISPR/CAS9, shRNA- mediated knock down techniques
- Excellent personal communication and organization skills (in German & English language)
- Extensive experience in the analysis of complex molecular data using NCBI and ENSEMBL and similar data bases.
- Broad knowledge with ADOBE/COREL DRAW graphic programs, statistical software and bioinformatics online tools (TRANSFAC, CLUSTAL, BLAST, etc).
- English (written and spoken) for publication work, participation in distinct student supervision work for Bachelor and Master research projects & research internships

## Contact & Application

Application should be submitted per mail or e-mail as one single PDF-file (max 5 MB) to Dr. Julia Rehnitz ([julia.rehnitz@med.uni-heidelberg.de](mailto:julia.rehnitz@med.uni-heidelberg.de)).

Please include an extensive motivation letter, your scientific CV, academic certifications including complete list of publications; 2-3 references of your past supervisors and give your expected availability date.

Please do not send original documents by mail services, since they will not be returned.

**For further information you may like to contact: [julia.rehnitz@med.uni-heidelberg.de](mailto:julia.rehnitz@med.uni-heidelberg.de)**



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