

FMR1 related premature ovarian insufficiency (OMIM# 311360)

Background

Premature ovarian insufficiency (POI) (also known as premature ovarian failure) is defined as the cessation of menses in women before the age of 40 for four or more months in association with FSH levels in the menopausal range.

FMR1 related POI is caused by a premutation-sized expansion of a CGG repeat within the 5'UTR of the *FMR1* gene (OMIM* 309550). Approximately 21% of female premutation carriers develop POI compared to only 1% of the general population.

FMR1 expansions in the premutation size range are likely to be unstable and prone to further expansion in the next generation. The identification of a premutation may therefore have implications for other family members who may inherit this expansion.

Recommended Clinical Referral Criteria

- **Diagnostic testing (Infertility referrals)**
Premature ovarian insufficiency
- **Family testing (Clinical Genetics referrals)**
Relatives with a confirmed family history of an *FMR1* related disorder

Molecular Analysis

- 1) **FRAXA PCR Assay:** Fragment length analysis of *FMR1* gene [CGG]_n triplet repeat. Pre-screen for the majority of diagnostic referrals.
- 2) **Asuragen Amplidex PCR Assay:** Fragment length analysis of *FMR1* gene [CGG]_n triplet repeat expansions in pre-mutation and full mutation size range by CGG repeat primed PCR analysis.

Prices* & Turnaround Times (TAT)

* Prices available on request

Test	TAT
<i>FMR1</i> expansion analysis (PCR)	42cd
<i>FMR1</i> high sensitivity gene expansion analysis	42cd

Contact Details

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Accredited to ISO 15189:2012
(8988)

Sample Requirements

Blood – 5ml in EDTA (1ml neonates/infants)
Please contact lab prior to sending a prenatal sample.
Please label samples with three identifiers and date of collection.

All samples must be accompanied by request form

Consent for testing & DNA storage is assumed when request for test received.

Links

Orphanet

<http://www.orpha.net/>

OMIM

<http://www.omim.org/>

Genetic Test Registry

<http://www.ncbi.nlm.nih.gov/gtr/>

Support

www.fragilex.org.uk