Mental Status of Females with an FMR1 Gene Full Mutation

B. B. A. de Vries,1 A. M. Wiegers,3 A. P. T. Smits,4 S. Mohkamsing,1 H. J. Duivenvoorden,2 J.-P. Frys,5 L. M. G. Curfs,3,* D. J. J. Halley,1 B. A. Oostra,1 A. M. W. van den Ouweland,1 and M. F. Niermeijer1

Departments of 1Clinical Genetics and 4Medical Psychology and Psychotherapy, University Hospital Dijkzigt and Erasmus University, Rotterdam; 2Research Department Observation Centre De Hondsberg, Oisterwijk; 3Department of Human Genetics, University Hospital Nijmegen, Nijmegen; and 5Division of Human Genetics, University Hospital Gasthuisberg, Leuven, Belgium

Summary

The cloning of the FMR1 gene enables molecular diagnosis in patients and in carriers (male and female) of this X-linked mental retardation disorder. Unlike most X-linked disorders, a considerable proportion of the male carriers of a full mutation in the FMR1 gene is affected. In this study, the intelligence quotients (IQs) were ascertained by the Wechsler Adult Intelligence Scale in 33 adult females with a full mutation, with 28 first-degree adult female relatives (mainly sisters) without a full mutation as controls. Seventy-one percent of the females with a full mutation had IQ scores below 85. In paired analysis, no significant correlation could be detected between the IQs of the females with a full mutation and those of their first-degree female relatives, reflecting a dominant effect of the FMR1 gene full mutation in the mental development of females. Considering females with a full mutation only, we observed a significant relation between the proportion of normal FMR1 alleles on the active X chromosome and IQ. We present a model to explain this relationship.

Introduction

The fragile X syndrome is the most common single cause of inherited mental retardation, affecting (according to earlier estimates) ±1/1,250 males and ±1/2,000 females (Webb et al. 1986). Recently, a lower incidence was reported of ±1/4,000 for males (Turner et al., in press). The cloning of the FMR1 gene enabled direct DNA diagnosis in both male and female fragile X patients (Fu et al. 1991; Oberlé et al. 1991; Verkerk et al. 1991; Yu et al. 1991). The FMR1 gene contains a variably sized CGG trinucleotide repeat at the 5' end, which has a length of between 6 and 53 repeats in normal individuals. Patients have >200 CGG repeats, the so-called full mutation, which is associated with methylation of a CpG island upstream of the FMR1 gene and with absence of gene transcription (Oberlé et al. 1991; Pieretti et al. 1991). Normal carrier males and females, with an exception for normal females with a full mutation, have an intermediate repeat number of between 43 and 200, the so-called premutation, with an unmethylated CpG island in front of the FMR1 gene and normal transcription (Pieretti et al. 1991). The FMR1 gene contains a variably sized CGG trinucleotide repeat at the 5' end, which has a length of between 6 and 53 repeats in normal individuals. Patients have >200 CGG repeats, the so-called full mutation, which is associated with methylation of a CpG island upstream of the FMR1 gene and with absence of gene transcription (Oberlé et al. 1991; Pieretti et al. 1991). Normal carrier males and females, with an exception for normal females with a full mutation, have an intermediate repeat number of between 43 and 200, the so-called premutation, with an unmethylated CpG island in front of the FMR1 gene and normal transcription (Pieretti et al. 1991).

The fragile X syndrome is transmitted as an X-linked semidominant disorder, since males and females may be affected. In earlier studies, the cognitive profile in females with the fragile X syndrome has been related to fragile X chromosome expression. Some groups found a significantly negative relationship between the percentage of fragile X expression and intelligence quotient (IQ) (Chudley et al. 1983; Wilhelm et al. 1988). Others found no such relation between IQ and fragility (taking ≥2% of fragile X cells as evidence for the carrier status) (Cronister et al. 1991).

Before the cloning of the gene, the reported incidence of intellectual deficits (IQ < 85) among heterozygous females ranged from 35% to 53% (Sherman et al. 1985; Hagerman et al. 1992). Besides the limitations of cytogenetic methods for the diagnosis of heterozygous females, most of those earlier studies also lacked an adequate control group. Recent studies based on DNA mutation analysis show mental impairment in 52%–82% of the females with a full mutation (Rousseau et al. 1991a, 1991b; Smits et al. 1994; Taylor et al. 1994), whereas Reiss et al. (1993) showed that the premutation in females did not affect their intellectual development. Molecular analysis of the FMR1 gene in carrier fe-
The percentages of normal FMR1 alleles on the active X chromosomes (NA) and normal FMR1 alleles on the inactive X chromosomes (NI) were estimated by determining the beginning and the end of the smear on Southern blot in carriers of the full mutation (fig. 1).

The size of the full mutation was estimated by averaging the smallest and largest detectable expansion. Sizes were estimated by determining the beginning and the end of the smear on Southern blot in carriers of the full mutation (fig. 1).

The percentages of normal FMR1 alleles on the active X chromosomes (NA) and normal FMR1 alleles on the inactive X chromosomes (NI) were ascertained through a male index patient (usually a first-degree relative) and were studied after informed consent was obtained. The study was approved by the Medical Ethical Committee of the University Hospital, Rotterdam. One female did not want to participate (not counted). Twenty-one females were paired to a sister with a normal FMR1 genotype (aged 22–72 years, mean 41.3 ± 14.5 years). In case more female control sibs were available, the female with the best age match was included in the study. In the absence of a control sister, the mother with a premutation was included as a control X chromosome (NI). For five females with a full mutation from large sibships, a unique control was not available, because every control had only been used once. Those five females were excluded from paired analysis (leaving 28 pairs) but were included in the molecular association studies.

**Determination of IQ Levels**

In all 61 females, the IQ and profile were ascertained using the Dutch version of the Wechsler Adult Intelligence Scale. (Stinissen et al. 1970). IQ testing was performed at the home of females by one examiner (A.W.) who was not informed about the genetic status of the women.

**DNA Analysis**

The intragenic DNA probe pP2 was used for DNA analysis of the FMR1 gene (Oostra et al. 1993). Genomic DNA was isolated from blood leukocytes (Miller et al. 1988). DNA (8 μg) was digested to completion with the restriction enzymes HindIII and Eagl according to the manufacturer’s instructions, separated by gel electrophoresis, and subjected to Southern blot analysis according to standard procedures (Sambrook et al. 1989). The probe was labeled by the random oligonucleotide priming method (Feinberg and Vogelstein 1983). After prehybridization and hybridization, the filters were washed to 0.1 × SSC at 65°C prior to autoradiography (Sambrook et al. 1989). Several autoradiograms (2–3) with different exposure times were made.

The size of the full mutation was estimated by averaging the smallest and largest detectable expansion. Sizes were estimated by determining the beginning and the end of the smear on Southern blot in carriers of the full mutation (fig. 1).

The percentages of normal FMR1 alleles on the active X chromosomes (NA) and normal FMR1 alleles on the inactive X chromosomes (NI) were ascertained by densitometry using a scanner (HP, scanjet IICX). The proportion of normal FMR1 alleles on the active X chromosome (NA) versus normal FMR1 alleles on the inactive X chromosome (NI) was calculated using the equation: NA/(NA + NI) (fig. 1) (Rousseau et al. 1991b).

In females with a mosaic pattern of the FMR1 gene mutation (full mutation with an additional premutation), estimation of the size of the expansion and the proportion of normal active X will be hampered by the additional premutation. Therefore, females with a premutation in addition to the full mutation (n = 6) were excluded from the comparison of the size of the full mutation and the X-inactivation ratio with IQ scores.

**Data Analysis**

A paired t-test for continuous data was applied to test for significant differences between the female carriers of the full mutation and their controls. The bivariate relationship between variables was estimated by means of Pearson’s product-moment correlation coefficient (r).
This correlation coefficient was considered to be an indicator of relative importance, while the corresponding P value represents the level of significance statistically. Adjusting for age, the partial correlation coefficient was the indicator of relative importance.

Results

IQ and Profile in Females with the Full Mutation versus Controls

The mean full-scale IQ (FSIQ), performance IQ (PIQ), and verbal IQ (VIQ) of the different groups studied are shown in table 1. The 28 female carriers of a full mutation had significantly lower mean FSIQ, PIQ, and VIQ scores than their paired controls with a normal FMR1 gene or with a premutation (for all means P < .001, paired-sample t-test). There was no significant difference of FSIQ between the females with a full mutation only (n = 27) and the females with a full mutation and an additional premutation, the so-called mosaics (n = 6) (r = .22; P = .21; table 1). Also, no significant difference of FSIQ, PIQ, and VIQ was observed between the females with a normal FMR1 genotype and the females with a premutation (r = -.1, P = .61; r = -.17, P = .39, respectively, r = -.02, P = .9).}

Paired Analysis: Females with a Full Mutation Compared with the Controls

In figure 2, each pair (female with a full mutation and control) and each female’s IQ are shown. In the group of 28 pairs, 20 (71%) of 28 carriers of the full mutation had an IQ <85 points. When the more strict criteria for mental retardation (IQ < 70 points) was used, 14 (50%) of 28 carriers of the full mutation were mentally retarded, whereas none of the control females had an IQ <70 points. Sixteen (57%) of 28 carriers of the full mutation scored >30 IQ points (2 SD) lower than their control. This ratio remained nearly unchanged when only the 21 sister-sister pairs are considered (62%). Remarkably, two females with the full mutation had a higher IQ than did their controls (fig. 2). No significant correlation could be detected between the FSIQ of the females with the full mutation and the FSIQ of the controls (r = .22; P = .27). Likewise, there was no significant correlation for the VIQ of the females with a full mutation and the VIQ of the controls (r = .07; P = .74), but the PIQ correlation was significant (r = .39, P = .04).

The Size of the FMR1 Gene Mutation and Mental Status in Females with the Full Mutation Only

For females with a full mutation, only the FSIQ, PIQ, and VIQ scores were negatively related with age (r...
Discussion

The Full Mutation in FMR1 Gene in Females Has a Dominant Effect over Other Factors Involved in Intellectual Development

In the present study, 71% (20/28) of the females with a full mutation in the FMR1 gene had IQ scores <85, representing borderline or mild/moderate retardation. Other groups also using DNA analysis of the FMR1 gene reported lower percentages, varying from 52% to 65%, of mental impairment (IQ < 85) among the females with a full mutation (Rousseau et al. 1991b; Taylor et al. 1994) (table 2), except for Smits et al. (1994) who reported 82%. However, in three of these studies, the mental development was clinically estimated without accurate IQ testing (Rousseau et al. 1991b; 1994; Smits et al. 1994).

Using strict criteria for mental retardation (i.e., IQ < 70 points), 50% (14/28) of the females with a full mutation in this study were mentally retarded. This is higher than the observation of Taylor et al. (1994), who reported 22% (5/23) of the females to have an IQ <70. Their results might be biased by the use of four different IQ tests and the low average age of the subjects they studied. Further, they could not exclude an ascertainment bias because of “unavoidable selection for motivation to come to the clinic” (p. 513), which is a major problem in several comparable studies. The present study avoided such a bias, both by selecting the families through a male index patient and by visiting the families at their homes. Although the possibility that females with a full mutation, ascertained through a male index patient, might be functioning on a different cognitive level as females with a full mutation without an affected, first-degree, male relative might be conceivable, at present no difference of this type is known.

The first-degree female relatives without a full mutation in the FMR1 gene (mainly sisters without an FMR1 gene...
### Table 2

**Summary of Molecular and IQ Data in Female Carriers of the Full Mutation: Overview of the Literature and the Present Study**

<table>
<thead>
<tr>
<th>Study</th>
<th>N</th>
<th>Age (years)</th>
<th>Controls</th>
<th>IQ Test</th>
<th>IQ &lt; 85 (%)</th>
<th>Size of FM&lt;sup&gt;a&lt;/sup&gt; to IQ</th>
<th>X inactivation&lt;sup&gt;b&lt;/sup&gt; to IQ</th>
<th>X inactivation&lt;sup&gt;b&lt;/sup&gt; to Age</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rousseau et al. (1991) (France)</td>
<td>27</td>
<td>All</td>
<td>Women PM&lt;sup&gt;c&lt;/sup&gt;</td>
<td>Clinical estimate</td>
<td>52</td>
<td>?</td>
<td>Yes (+)</td>
<td>Yes (+)</td>
</tr>
<tr>
<td>Taylor et al. (1994) (Denver, USA)</td>
<td>23</td>
<td>3–58</td>
<td>Mothers</td>
<td>Test (4)&lt;sup&gt;d&lt;/sup&gt;</td>
<td>65</td>
<td>No</td>
<td>No</td>
<td>Yes (+)</td>
</tr>
<tr>
<td>Abrams et al. (1994)&lt;sup&gt;e&lt;/sup&gt;</td>
<td>31</td>
<td>4–27</td>
<td>Mothers (+fathers)</td>
<td>Test (2)&lt;sup&gt;d&lt;/sup&gt;</td>
<td>?</td>
<td>Yes (-)</td>
<td>Yes (+)</td>
<td>No</td>
</tr>
<tr>
<td>Reiss et al. (1995)&lt;sup&gt;e&lt;/sup&gt; (Baltimore, USA)</td>
<td>29</td>
<td>6–17</td>
<td>Mothers + fathers</td>
<td>Wechsler Intelligence Scale for Children</td>
<td>?</td>
<td>No</td>
<td>Yes (+)</td>
<td>?</td>
</tr>
<tr>
<td>Rousseau et al. (1994) (collaborative)</td>
<td>170</td>
<td>All</td>
<td>Women N&lt;sup&gt;f&lt;/sup&gt; and PM&lt;sup&gt;e&lt;/sup&gt;</td>
<td>Clinical estimate</td>
<td>59</td>
<td>Yes (-)</td>
<td>No</td>
<td>Yes (+)</td>
</tr>
<tr>
<td>Present study</td>
<td>33</td>
<td>20–70</td>
<td>Sisters (+mothers)</td>
<td>Weschler Adult Intelligence Scale</td>
<td>71</td>
<td>No</td>
<td>Yes (+)</td>
<td>No&lt;sup&gt;g&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

**NOTE.**—Minus sign (-) or plus sign (+) correspond to negative or positive correlation, respectively.

<sup>a</sup> Full-mutation FMR1 gene.

<sup>b</sup> Proportion of normal active X, as described by Rousseau et al. (1991b).

<sup>c</sup> Premutation FMR1 gene.

<sup>d</sup> Number of different IQ tests used.

<sup>e</sup> Study groups partly overlap.

<sup>f</sup> Normal FMR1 gene.

<sup>g</sup> Adults only.

The full mutation in the FMR1 gene in females can be characterized by different features, such as size of the mutation and localization on the active or inactive X chromosome. No relation between the size of the full FMR1 mutation, whether characterized by smallest, average, or largest size, and the IQ level of the females was observed in this study. These findings are inconsistent with earlier reports by Abrams et al. (1994) and Rousseau et al. (1994), but they are in line with the reports from other groups (Rousseau et al. 1991b; Taylor et al. 1994; Reiss et al. 1995) (table 2). Because a relation between size and IQ level could not be detected in males (De Vries et al. 1993; Loesch et al. 1993), it would be unlikely to observe such relationship in females. This corresponds with the characteristics of the full mutation in the FMR1 gene. Independent of the size of a full mutation, the full mutation is associated with methylation of a CpG island in front of the FMR1 gene, leading to an absence of FMR1 protein.

As in other X-linked disorders, lyonization may influence the phenotypic expression of the normal allele in females with a full mutation in the FMR1 gene. Earlier studies using cytogenetic techniques in lymphocytes explored whether the normal X chromosome is more frequently the active (or early replicating) X chromosome in carrier females with a normal intelligence compared to carriers with an intellectual impairment. Although some studies confirmed the relation between a high percentage of normal active X chromosomes and normal intelligence (Knoll et al. 1984; Paul et al. 1984; Wilhelm et al. 1988), others did not (Nielsen et al. 1983; Fryns et al. 1984). With molecular techniques, the proportion of normal FMR1 alleles on the active X chromosome...
can be estimated more accurately by using methylation assays. The cells with normal active FMR1 alleles are actually capable of producing FMR1 protein. In the 27 females with a full mutation only, the normal FMR1 alleles were significantly more frequent on the active X chromosome in intellectually higher-functioning females than in females with a mental impairment. This confirms earlier findings (Abrams et al. 1994; Rousseau et al. 1991b; Reiss et al. 1995) but contradicts two other studies (Rousseau et al. 1994; Taylor et al. 1994) (table 2). In the study by Rousseau et al. (1994), the intelligence level was clinically estimated without specific IQ testing, and therefore the relation might have been missed. Taylor et al. (1994) included females with a mosaic pattern (premutation and full mutation). Although we did not observe a difference of the FSIQ between females with a full mutation versus mosaics, the effects of an additional premutation on the phenotype have not been fully assessed in larger controlled studies. Therefore, including "mosaic" females might influence the results. Another reason for excluding "mosaic" females is the inability to establish the percentage of normal active X chromosomes accurately in these cases.

We observed an age-independent, skewed X inactivation in leukocytes favoring the X-chromosomes with normal FMR1 alleles over those with a full-mutation FMR1 allele in females both with normal and subnormal intelligence level. This suggests a positive selection for leukocytes able to produce FMR1 protein earlier in life. Some groups (Rousseau et al. 1991b, 1994; Taylor et al. 1994) observed an age-dependent selection in leukocytes: a higher proportion of active X chromosomes carrying the normal FMR1 gene in elderly women with a full mutation compared to younger females. Because only adults were included in the present study, we could not observe such an age-dependent selection.

In each cell, the X inactivation occurs at random in the late blastocyst stage (Nesbitt 1971; McMahon et al. 1983; Lyon 1994). The precursor cells for both brain and blood formation are the embryonic ectodermal cells, which undergo X inactivation around the 2d wk of embryonic development. The X-inactivation distribution is rather consistently equal when different tissues within one individual are compared, both in mice and men (Nesbitt 1971; Fialkow 1973, McMahon et al. 1983; Kolehmainen and Karant 1994). Because we observed females with a high proportion of normal FMR1 alleles (>.80) with either a normal or a subnormal IQ level, the currently observed relation is likely influenced by the age-dependent selection for normal FMR1 expression in leukocytes. We hypothesize that the proportion of inactivation of the normal X chromosome at the late blastocyst stage in some individuals is <.5 and in others >.5. Individuals with a lower proportion of normal active X chromosome in the brain will have subnormal intelligence, while individuals with a higher proportion will be normal. There will be a relation between the proportion of normal FMR1 alleles on the active X chromosome in the brain and IQ. In leukocytes, however, there is selection for normal X chromosomes, leading to a change in the proportion of normal X chromosomes. This change is more likely to be higher in individuals with a low proportion of normal X chromosomes, disturbing the relationship between proportion of normal alleles and IQ. The presently observed relation between a higher proportion of normal FMR1 alleles on the active X chromosomes in leukocytes and a higher intelligence in females with a full mutation might still be a reflection of the distribution of X inactivation in brain cells. To prove this hypothesis, we propose to study the X-inactivation pattern in other tissues that might lack this selection phenomenon. Studies on protein expression in neuronal cells of either females with a full mutation or heterozygous knock-out mice might give the relevant clues.

The currently observed relation between molecular parameters and the FSIQ were also significant for the subcomponent PIQ but not for VIQ. This is in line with a recent report about girls with a full mutation, in which a stronger relation between the proportion of normal active X was suggested for PIQ than for VIQ (Reiss et al. 1995). Since the PIQ is less influenced by cultural factors than is the VIQ (Cattel 1963; Horn 1968), the PIQ seems preferable over the VIQ and FSIQ as an indicator of mental developmental capabilities in molecular-cognitive association studies.

The mental status in females with the full mutation is influenced by several factors. Besides the full mutation in the FMR1 gene, other genetic and environmental factors may play, to an unknown extent, a role in intellectual development. In this first study of comparing full mutation FMR1 female carriers with their noncarrier sisters, those shared factors were less significant, reflecting a dominant effect of the FMR1 gene full mutation on mental development in a majority of females with a full mutation. This finding will be relevant in the genetic counseling of carriers of (pre-)mutations of the fragile X syndrome. X-inactivation studies during prenatal diagnosis in female pregnancies with a full FMR1 mutation will not allow a reliable prognosis of the intellectual development of such a child. Henceforth, parents will have to decide on the general risk estimates for intellectual impairment in female pregnancies with a full mutation as observed in this study.

Acknowledgements

We are thankful to Prof. Dr. H. Galjaard and the Foundation for Clinical Genetics, Rotterdam, for their continuous
support, and Prof. Dr. B. A. van Oost for providing some of the DNA samples and his comments on the manuscript.

References


Sutherland GR (1977) Fragile sites on human chromosomes: demonstration of their dependence on the type of tissue culture medium. Science 197:265–266