Short reports

A family with mental retardation, variable macrocephaly and macro-orchidism, and linkage to Xq12-q21

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Abstract

A family with X linked inheritance of mental retardation (XLMR) is presented. There are 10 mentally retarded males and two affected females in two generations. There are four obligatory carriers, one of whom is described as "slow". Most affected males show macrocephaly and macro-orchidism, which are typical signs of the fragile X syndrome, but have been tested cytogenetically and by analysis of the FMR1 gene and do not have this syndrome. However, some normal males in the family also exhibit macro-orchidism and macrocephaly. Linkage analysis using markers derived from the X chromosome indicates that the causative gene in this family is located in the proximal long arm of the X chromosome, in the interval Xp11-q21. Maximum lod scores of 2.96 with no recombination were found at three loci in Xq13-q21: DXS1111, DXS566, and DXS986. Recombination was observed with DXS1002 (Xq21.31) and DXS991 (Xp11.2), loci separated by about 30 Mb. Although isolation of the gene in this family will be difficult because of the size of the region involved, the localisation should be helpful in investigating other similar families with XLMR, macrocephaly, and macro-orchidism not attributable to FMR1.

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Non-specific X linked mental retardation (NSXLMR) is a clinical entity presenting difficulties in categorisation because of lack (by definition) of an associated phenotype other than mental retardation.^{1,2} Linkage data are now available for over 40 NSXLMR families, with unique genes (denoted as MRX with unique numbers) potentially responsible for the mental retardation in each family.³ However, overlapping linkage results indicate that only a few of these genes map to exclusive locations.^{4,5}

Early in the development of the clinical definition of fragile X syndrome, there was recognition that macro-orchidism, one of the cardinal associated findings, could also occur in NSXLMR males not manifesting X chromosome fragility.⁶⁻⁸ In addition, macrocephaly is a common finding in XLMR.⁹ There are more recent reports of similarly affected subjects and families.¹⁰⁻¹² The terms fragile X negative Martin-Bell syndrome,¹³ mental retardation-macro-orchidism, X linked (MRMO¹⁴), or XLMR+MO¹⁵ (McKusick MIM No 309530) have been used to describe these families.

We describe a family with macrocephaly and macro-orchidism associated with XLMR in which some retarded males do not have both macrocephaly and macro-orchidism, but some unaffected males do. We present linkage data indicating that the causative gene is in the Xp11-q21 region.

Subjects and methods

FAMILY

This family was ascertained when subjects III.15, III.18, and III.20 were evaluated because III.13 wished to know her risk of having an affected son (fig 1). There are 10 mentally retarded males in two generations of this family, with two retarded females and 10 unaffected males (one male of uncertain phenotype died at 3 months of pneumonia). The inheritance of the mental retardation is consistent with X linked mental retardation (XLMR). Of interest, but presumably unrelated, is the fact that II.5 had neurofibromatosis (NF) and died of an islet cell carcinoma of the pancreas. None of the people we examined personally showed any signs of NF and this includes five of the affected males.

Clinical information for six of the affected and five of the unaffected males was available. Findings for these males and for five at risk females are summarised in table 1. Fragile X cytogenetic studies on the three index males were negative on two occasions in two different laboratories. FMR1 studies (not shown) by polymerase chain reaction and Southern blotting showed normal results on affected males. Intelligence was assessed by examination, report, or by IQ score when available. The IQ

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Figure 1 Partial pedigree of XLMR family. Only critical subjects are shown and not necessarily in order of age. Haplotypes for the indicated DNA markers are shown beneath pedigree symbols. Boxed portions of the haplotype segregate with mental retardation in affected males.

scores available for four of the affected males are remarkably consistent and fall within the range of moderate mental retardation. Four of the affected males have macro-orchidism with testicular volumes >35 ml. Volumes were estimated using a published formula.⁷ One mentally retarded male (III.15) has testes of normal volume (23 ml, normal upper limit 25–35 ml), and one was not examined. Surprisingly, two of the unaffected males (II.3 and III.19) also have testicular volumes >35 ml. Five of six examined affected males also have macrocephaly

with occipitofrontal circumferences (OFC) of >90th centile. The same two unaffected males (II.3 and III.19) have similar OFCs, as does an at risk female, III.13. Two presumably affected females are described as “slow” and are living with their parents as adults (III.3 and III.4). The carrier mother (II.4) is also described as “slow” (this branch of the family was unavailable for detailed study, as was the branch from II.6). The affected males have monotonous but not repetitive or “jocular” speech, blue eyes with dark hair, and a triangular facial appearance with some prominence of the jaw (fig 2), but no facial elongation or lateral prominence of the ears, which would be more reminiscent of the fragile X syndrome. Two affected males (II.8 and III.15) have a seizure disorder, as does an unaffected male, IV.1. The personality is outgoing without autistic characteristics, there is normal eye contact, and there are no hand flapping movements or self-abuse.

DNA ANALYSIS

The linkage project was approved by appropriate reviewers (see Acknowledgments) and informed consent was obtained from sampled people. DNA was isolated from leucocytes in standard fashion. Polymerase chain reaction (PCR) was accomplished using a Perkin Elmer 9600 thermocycler. The X chromosome markers spanning Xp22.31 to Xq28, for which

Table 1 Clinical data

No	Age	Sex	OFC (cm)	OFC (%)	Height (in)	Testes (ml)	IQ
II.1	40s	F				NA	NL
II.2	69	F				NA	NL
II.3	72	M	59	>98	67.5	98	NL
II.8	58	M	59	>98		58	MR
II.9	56	M	58.5	98		49	MR
III.1	39	M			71	32	NL
III.10	33	M			66	32	NL
III.13	29	F	57.5	98		NA	NL
III.14	27	M	56 at 6 y	>98			35
III.15	26	M	58	90	71	23	43
III.16	23	F	56	75		NA	NL
III.17	25	F	54	25		NA	NL
III.18	22	M	54	25		54	43
III.19	18	M	58.5	98	67.5	38	NL
III.20	17	M	61	>98		38	38
IV.1	7	M	53	75	47	2	NL

OFC: occipitofrontal circumference, %: centile, in: inches, cm: centimetres, IQ: intelligence quotient, NA: not applicable, NL: normal intelligence, MR: mentally retarded. Blanks indicate data are not available for the particular measure.

Figure 2 Affected males as indicated on the pedigree (fig 1). (Photographs reproduced with permission.)

inheritance in this family was analysed, are listed in table 2. Primers for the microsatellite markers were FAM labelled on a Beckman Oligo synthesiser using primer sequences from the Genome Data Base (GDB, Baltimore).

The total volume for each PCR reaction was 20 μ l. The reactions contained 100 ng DNA, 1 μ mol/l of each primer, 1 $\times$ Mg buffer, 0.2 mmol/l dNTPs, and 1/10 unit of *Taq* polymerase. Ten μ l of the PCR product from each reaction were mixed with 2 μ l of bromophenol blue

dye and loaded on 2% agarose gels in 1 $\times$ TBE buffer. Following electrophoresis to visualise results, the products were diluted and loaded onto an ALF (Pharmacia) gel for size analysis.

Allelic inheritance was interpreted with reference to CEPH control 133402, family members, marker standards, and published data. Linkage of the mental status phenotype (with "affected" being mentally retarded, regardless of associated macrocephaly or macro-orchidism) to alleles for the DNA

Table 2 X chromosome markers and linkage results

Marker	Locus	Location	Recombination							θ_{max}	Z_{max}
AFM120Xa9	DXS987	Xp22.31	0.001	0.01	0.05	0.1	0.2	0.3			
AFM184xg5	DXS992	Xp21.3	-11.82	-6.84	-3.45	-2.08	-0.86	-0.31	0.50	0.00	
AFM151xf6	DXS991	Xp11.21	-9.04	-5.07	-2.39	-1.34	-0.46	-0.10	0.43	0.03	
ARA	ARA	Xq12	-3.04	-1.08	0.17	0.57	0.72	0.58	0.19	0.73	
AFM078za1	DXS983	Xq12	2.95	2.91	2.72	2.47	1.93	1.31	0.00	2.96	
DXS1111	DXS1111	Xq12	0.67	0.66	0.61	0.55	0.40	0.23	0.00	0.67	
HX60	DXS566	Xq13.3	2.95	2.91	2.70	2.44	1.87	1.25	0.00	2.96	
AFM116xg1	DXS986	Xq21.1	2.95	2.91	2.71	2.45	1.89	1.27	0.00	2.96	
DXS738	DXS738	Xq21.1	2.95	2.91	2.72	2.45	1.90	1.28	0.00	2.96	
AFM207zg5	DXS995	Xq21.1	2.95	2.91	2.72	2.47	1.92	1.30	0.00	2.96	
AFM6357Xe5	DXS8076	Xq21.1	0.85	0.83	0.77	0.69	0.50	0.29	0.00	0.85	
AFM249vh5	DXS1002	Xq21.31	2.94	2.91	2.72	2.47	1.93	1.31	0.00	2.96	
AFM136yc7	DXS990	Xq21.33	-0.03	-0.03	-0.02	-0.10	0.00	0.00	0.39	0.00	
XL5A	DXS424	Xq23	-0.05	0.92	1.43	1.51	1.32	0.97	0.09	1.51	
AFM248we5	DXS1001	Xq24	-4.93	-2.93	-1.56	-0.98	-0.46	-0.19	0.50	0.00	
46	DXS548	Xq27.3	-14.74	-8.76	-4.67	-2.98	-1.44	-0.67	0.50	0.06	
sDF-2	DXS1108	Xq28	-7.88	-4.90	-2.92	-2.14	-1.40	-0.86	0.50	0.60	
			-15.04	-9.06	-4.97	-3.28	-1.72	-0.89	0.50	0.52	

markers was assessed by the FASTLINK modification of the LINKAGE program.^{16 17} Penetrance was assigned at 1.0 for mental retardation associated with the affected allele. Assignment of penetrance for females is a moot issue, as none of the four females genotyped inherited the "affected" haplotype.

Results

Linkage analysis results are shown in table 2. Three loci (DXS1111, DXS566, and DXS986) show linkage to the mental retardation phenotype in males at a maximum lod score of 2.96 with no recombination. Given the known map positions for these loci, the gene causing NSXLMR in this family is localised between Xp11 and Xq21. The region is flanked by the loci DXS991 in Xp11.2 and DXS1002 in Xq21.31, which show recombination and are separated by about 30 Mb. Alleles for the markers are shown in inferred haplotypes on the pedigree (fig 1). For loci DXS983, DXS1111, DXS566, DXS986, DSX738, and DXS995, respectively, the haplotype 2-4-3-1-2-2 is observed in all affected males. The at risk female with macrocephaly, III.13, does not share this haplotype, indicating that her macrocephaly is probably not a manifestation of carrier status. Similarly, III.17, who has a son with seizures, does not carry this haplotype, indicating that her son's seizures are probably unrelated to the XLMR allele in the family.

Discussion

The gene causing XLMR in this family is most probably located in the proximal Xq region. This positioning shows considerable overlap with results from other families with XLMR, including MRX 1, 4, 5, 6, 7, 8, 9, 13, 14, 17, 20, 22, 26, 30, and 31.³⁻⁵ In addition, specific XLMR genes for Menkes disease,¹⁸ PGK deficiency,¹⁹ and α thalassaemia/mental retardation (ATR-X),²⁰ and for other syndromes²¹⁻²⁹ map to this region.³ These latter syndromes are generally excluded in this family by lack of consistent clinical findings. Of the two recent reports of families with macro-orchidism, the family investigated by Hu *et al.*¹¹ with the causative gene designated MRX2, maps to Xp21-22, excluding overlap with the family in our report. Affected subjects in this previously reported family are similar to the males described here, with macrocephaly and macro-orchidism, but also with prominent ears. There is no map information for the family described by Tariverdian *et al.*¹⁰ However, this seems to be a different condition with males manifesting prominent eyebrows and jaws and abnormal ears in addition to macro-orchidism. Mutations in the ATR-X gene can produce multiple phenotypes, and therefore this is a "candidate" causative gene for the family described in our report. Potential identity with one of the other MRX genes cannot be excluded. Until the causative genes are isolated, correct designation of the location and number of XLMR genes cannot be defined.

In this family, although the physical phenotype in affected relatives somewhat resembles that in patients with the fragile X syndrome,

the characteristics contributing to this impression are inherited irregularly, with some unaffected males exhibiting macro-orchidism or macrocephaly or both, and one at risk female (with a non-carrier haplotype) showing macrocephaly. One explanation is that macro-orchidism and macrocephaly segregate coincidentally in this family. A similar family with macrocephaly, XLMR, and mild retardation in heterozygous females has been described by Turner *et al.*⁹ with linkage to the pericentromeric region. This location overlaps the Xp11-q21 location of the family described here. Isolation of the causative gene and investigation of other similar families will assist in correct categorisation of the XLMR conditions in this and other similar families.

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- 1 Turner G, Turner B. X-linked mental retardation. *J Med Genet* 1974;11:109-13.
- 2 Herbst DS. Nonspecific X-linked mental retardation 1: a review with information from 24 new families. *Am J Med Genet* 1980;7:443-60.
- 3 Lubs HA, Chiurazzi P, Arena JF, Schwartz C, Tranebjaerg, Neri G. XLMR genes: update 1996. *Am J Med Genet* 1996; 64:147-57.
- 4 Gedeon AK, Donnelly AJ, Mulley JC, Kerr B, Turner G. How many X-linked genes for non-specific mental retardation (MRX) are there? *Am J Med Genet* 1996;64:158-62.
- 5 Hane B, Schroer RJ, Arena JF, Lubs HA, Schwartz CE, Stevenson RE. Nonsyndromic X-linked mental retardation: review and mapping of MRX29 to Xp21. *Clin Genet* 1996; 50:176-83.
- 6 Jennings M, Hall JG, Hoehn H. Significance of phenotypic and chromosomal abnormalities in X-linked mental retardation (Martin-Bell or Renpenning syndrome). *Am J Med Genet* 1980;7:417-32.
- 7 Nielson KB, Tommerup N, Dyggve HV, Schou C. Macroorchidism and fragile X in mentally retarded males. *Hum Genet* 1982;61:113-17.
- 8 Fishburn J, Turner G, Daniel A, Brookwell R. The diagnosis and frequency of X-linked conditions in a cohort of moderately retarded males with affected brothers. *Am J Med Genet* 1983;14:713-24.
- 9 Turner G, Gedeon A, Mulley J. X-linked mental retardation with heterozygous expression and macrocephaly: pericentromeric gene localization. *Am J Med Genet* 1994;51:575-80.
- 10 Tariverdian G, Froster-Iskenius U, Deuschl G, Wolff G. Mental retardation, acromegalic face and megalotestes in two half-brothers: a specific form of X-linked mental retardation without fra(X)? *Am J Med Genet* 1991;38:208-11.
- 11 Hu LJ, Blumenfeld-Heyberger S, Hanauer A, Weissenbach J, Mandel J-L. Non-specific X-linked mental retardation: linkage analysis in MRX2 and MRX4 families revisited. *Am J Med Genet* 1994;51:569-74.
- 12 Chiurazzi P, de Graaff E, Ng J, *et al.* No apparent involvement of the FMR1 gene in five patients with phenotypic manifestations of the fragile X syndrome. *Am J Med Genet* 1994;51:309-14.
- 13 Opitz JM, Sutherland GR. International workshop on the fragile X and X-linked mental retardation. *Am J Med Genet* 1984;17:5-94.
- 14 Howard-Peebles PN, Finley WH. Screening of mentally retarded males for macro-orchidism and the fragile X chromosome. *Am J Med Genet* 1983;15:631-5.
- 15 McKusick VA. *Mendelian inheritance in man: catalogs of autosomal dominant, autosomal recessive, and X-linked phenotypes*. Baltimore: The Johns Hopkins University Press, 1994.
- 16 Cottingham RW Jr, Idury RM, Schaffer AA. Faster sequential genetic linkage computations. *Am J Hum Genet* 1993;53:226-35.
- 17 Lathrop GM, Lalouel JM. Easy calculation of lod scores and genetic risks on small computers. *Am J Hum Genet* 1984;36:460-5.
- 18 Horn N, Stene J, Mollekaer AM, Friedrich U. Linkage studies in Menkes disease: the Xg blood group system and C-banding of the X chromosome. *Ann Hum Genet* 1984;48:161-72.
- 19 Sugie H, Sugie Y, Nishida M, *et al.* Recurrent myoglobinuria in a child with mental retardation: phosphoglycerate kinase deficiency. *J Child Neurol* 1989;4:95-9.

- 20 Gibbons RJ, Picketts DJ, Villard L, Higgs DR. Mutations in a putative global transcriptional regulator cause X-linked mental retardation with thalassemia. *Cell* 1995;80:837-45.
- 21 Porteous MEM, Johnson H, Burn J, *et al.* A new mental retardation syndrome mapping to the pericentromeric region of the X-chromosome. *Am J Hum Genet* 1991;51:A106.
- 22 Fishbeck KH, ar-rushdi N, Rozear M, Pericak-Vance M, Fryns J. X-linked neuropathy: gene localization with DNA probes. *Am J Hum Genet* 1985;37:A153.
- 23 Goldblatt J, Ballo R, Sachs B, Moosa A. X-linked spastic paraplegia: evidence for homogeneity with a variable phenotype. *Clin Genet* 1989;35:116-20.
- 24 Schwartz CE, Ullmer J, Brown A, Pancoast I, Goodman HO, Stevenson RE. Allan-Herndon syndrome II. Linkage to DNA markers in Xq21. *Am J Hum Genet* 1990;47:454-8.
- 25 Wieacker P, Wolff G, Wienker TF. Close linkage of the Wieacker-Wolff syndrome to the DNA segment DXYS1 in proximal Xq. *Am J Med Genet* 1987;28:245-53.
- 26 Schwartz CE, Ouzts L, Gibson A, *et al.* Renpenning syndrome: evidence for pericentric location of the gene in two families, including the original Renpenning family. Paper presented at the 6th International Workshop of Fragile X and X-linked Mental Retardation, Palm Cove, Australia, August 3-6, 1993.
- 27 Gedeon A, Mulley J, Haan E. Gene localisation for Sutherland-Haas syndrome (SHS: MIM 309470). *Am J Med Genet* 1996;64:78-9.
- 28 Gedeon A, Mulley J, Turner G. Gene localisation for Wilson-Turner syndrome (WTS: MIM 309585). *Am J Med Genet* 1996;64:80-1.
- 29 Miles JH, Carpenter NJ. Unique X-linked mental retardation syndrome with fingertip arches and contractures linked to Xq21.31. *Am J Med Genet* 1991;38:215-23.