Small Deletion at the 7q21.2 Locus in a CCM Family Detected by Real-Time Quantitative PCR

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Cerebral cavernous malformations (CCMs) represent a common autosomal dominant disorder that predisposes patients to haemorrhagic strokes and focal neurological signs. About 56% of the hereditary forms of CCMs have been so far associated with mutations in the KRIT1 (Krev Interaction Trapped 1) gene, located at 7q21.2 (CCM1 locus). We described the complete loss of 7q21.2 locus encompassing the KRIT1 gene and 4 flanking genes in a CCM family by using a dense set of 12 microsatellite markers. The complete loss of the maternal copy of KRIT1 gene region was confirmed by Real-Time Quantitative Polymerase Chain Reaction (RT-QPCR) and the same approach was used for expression analysis. Additional RT-QPCR analysis showed the extension of the deletion, for a total of 700 kb, to the adjacent downstream and upstream-located genes, MTERF, AKAP9, CYP51A1, as well as a partial loss of theANKIB1 gene. Here we report the molecular characterization of an interstitial small genomic deletion of the 7q21.2 region in a CCMs affected family, encompassing the KRIT1 gene. Our findings confirm the loss of function mechanism for the already known CCM1 locus, without any evident involvement of the other deleted genes. Moreover, our investigations highlight the usefulness of the RT-QPCR to the molecular characterization of the breakpoints genomic deletions and to the identification of internal deleted genes involved in the human genetic diseases.

1. Introduction

Cerebral cavernous malformations (CCMs, [MIM 116860]) are congenital vascular defects mostly located within the CNS with a prevalence of 0.1%–0.5% in the general population [1]. They are characterized by abnormally enlarged capillary cavities without intervening brain parenchyma [2]. The presence of these altered vascular structures is believed to account for all symptoms, ranging from headache to focal neurological defects and rarely acute bleeding. CCMs are also associated to an increased probability of stroke and epilepsy [1, 3, 4]. Family cases are often characterized by the presence of multiple lesions with an incomplete penetrance probably related to gene involved [5, 6] and on the age at onset. Heritable CCMs have been so far associated with mutations in three genes: the KRIT1 (Krev Interaction Trapped 1) [7] and the CCM2 (Cerebral Cavernous Malformation 2, Malcavernin) genes located, respectively, at the 7q21.2 (CCM1 locus) and 7p15-p13 (CCM2 locus) [8], and the PDCD10 (Programmed Cell Death 10) gene is located at the 3q26.1-27 (CCM3 locus) [9]. KRIT1 gene is responsible for about 56% of the hereditary forms of CCMs, whereas the MGC4607 gene accounted for 33% of them. The third locus identified by mutational screening of the PDCD10 gene
2. Patients and Methods

2.1. Patients. All the subjects belonged to a Northern Italian family affected by cerebral cavernous malformation. The index case is a 36-year-old male with a seizure history and cerebral haemorrhages. The five family members were investigated by brain TC and MRI and the diagnosis of CCM was based on its characteristic radiographic findings. Each subject underwent detailed clinical assessment, with emphasis on neurological, dermatological, and ophthalmological examinations. Informed consent was obtained from all family members to perform genetic analyses.

2.2. Microsatellite Analysis. DNA was extracted from peripheral blood using a standard phenol-chloroform protocol [16]. Linkage analysis was performed in order to identify the disease associated gene. Haplotype analyses were performed for all CCMs family members by using a dense set of 13 microsatellite markers flanking the CCM1 locus (See Table 1 in Supplementary Material available online at doi:10.1155/2010/854737). Physical distances between markers were based upon the electronic database available from the University of California Santa Cruz (http://www.genome.ucsc.edu/). Amplifications were carried out in 25 μl reaction volume containing 100 ng of DNA, 10X PCR Buffer with 15 mM MgCl2, 200 μM each dNTPs, 20 pmol each primer, and 1 U AmpliTaq Gold DNA polymerase (Applied Biosystems, Foster City, CA). PCR cycling conditions consisted of initial 12 minutes denaturation step at 95°C, followed by 35 cycles of 95°C for 30 seconds, annealing for 30 seconds, and extension at 72°C for 30 seconds, with final extension at 72°C for 7 minutes. PCR products were visualised by ethidium bromide staining on 2% agarose gel. PCR products were loaded on a capillary electrophoresis ABI 3100 (Applied Biosystems), and results were analysed by using Genescan 3.7 and Genotyper 3.7 NT software (Applied Biosystems).

2.3. KRIT1 Gene Screening. The mutational analysis of the KRIT1 gene was performed as previously described [17]. Briefly, amplifications for all the 16 KRIT1 coding regions, including the exon-intron boundaries, were carried out and analysed by denaturing high-performance liquid chromatography (DHPLC; Transgenomic Inc. Transgenomic, Inc. Nebraska, USA) screening (primers sequences, annealing temperatures, and size of PCR products are in Supplementary Table 1; oven temperatures and acetonitrile gradients are available from the authors). Amplicons with an abnormal elution profile were purified using the GFX PCR and Band Purification Kit (Ge HealthCare, Buckinghamshire, UK), sequenced with the BigDye Terminator Cycle Sequencing Kit v. 1.1 (Applied Biosystems), loaded on ABI 3100 capillaries (Applied Biosystems) and analysed using the Sequencing Analysis software v2.0.

2.4. RT-QPCR Gene Copy Number Analysis. A new protocol of RT-QPCR was developed to provide a sensitive method for detecting large deletions encompassing the KRIT1 gene and the 7q21 locus.

Three sets of primers were designed for amplification of 7q21 region around the KRIT1 gene (Supplementary Table 2). All primers were designed with Primer Express 2.0 software (Applied Biosystems) and tested for specificity using BLAT software (http://genome.ucsc.edu/cgi-bin/hgBlat?command=start). Primer sets 1 identify 8 amplicons covering the KRIT1 gene (exons 3, 9, 17, and 20) and 5’ and 3’ flanking 4.5 Kb regions. Primer set 2 annealed on the 10 KRIT1 flanking genes located into the 7q21 region delimited by D7S2410 and D7S646 microlistaties: PFTK1 (NM_012395), FZD1 (NM_003505), MTERF (NM_006980), AKAP9 (NM_005751), CYP51A1 (NM_000786), ANKBI1 (NM_019004.1), GATA1 (NM_021167), ERVWE1 (NM_014590), PEX1 (NM_000466), and CDK6 (NM_001259).

The identification of ANKBI1 gene breakpoint region was performed using an additional set of primers (set 3) designed on exons 7–13 of the gene (Supplementary Table 3). All primers were purchased by PRIMM (PRIMM Labs, Inc. MI, Italy).

Deletion of KRIT1 gene and of the flanking genes was detected on the 384-well ABI Prism 7900 Sequence Detection System (Applied Biosystems) by the measurements of the amplicons Copy Number (CN) using a RT-QPCR approach with SYBR-Green I detection. Reaction mixture (10 μl) contained 2.5x Fast Start DNA master mix hybridization SYBR Green (Roche Molecular Biochemicals, Mannheim, Germany), 250 μM of each forward and reverse primers, and 30 ng of DNA as template. Reactions were run on ABI PRISM 7900HT Sequence Detection System (Applied Biosystems). Cycling conditions were as follows: 10 minutes at 95°C, 40 cycles at 95°C for 15 seconds, and 60°C for 60 seconds. After PCR amplification, a melting curve was generated for every PCR product in order to verify the specificity of the PCR reaction. Calculation of the gene copy number was made using the 2−ΔΔCT method as reported (User Bulletin #2, Applied Biosystems) [18, 19]. Outlier values with a differences between Ct and Ct mean >0.3 were excluded from further data analysis. For normalization of the
relative amount, the gene copy numbers were divided by the geometric mean of two described reference with a normal copy number, ZNF80 (3q13.31) and MOX2 (3q13.2) [20]. Using this method, a Diploid Copy Number (D-CN) of 1.0 ± 0.2 is expected for a normal sample and a value of 0.5 ± 0.2 for a sample with 7q21 genomic deletion (Haploid Copy Number, H-CN).

2.5. Sample Processing, RNA Extraction and cDNA Synthesis. Total RNA of I : 1, I : 2, II : 1, and II : 3 subjects was extracted using PAXgene Blood RNA kit (PreAnalytix, Qiagen, Germantown, MD). RNA was eluted in RNAse free-water and stored at −80°C until used. RNA quality and concentration were measured by using 2100 Expert Analyzer (Agilent Technology, Inc. Headquarters) with an RIN (RNA Integrity Number) ≥9.0. After heating at 65°C for 5 minutes in order to denature RNA and to inactivate RNases, 500 ng of total RNA was subjected to reverse transcription using the QuantiTect Reverse Transcription Kit (Qiagen). cDNA was synthesized according to the manufacturer’s instructions.

2.6. RT-QPCR Gene Expression Analysis. Primers set were designed with Primer Express 2.0 software (Applied Biosystems) across the coding region of flanking KRIT1 genes into the 7q21 deleted region: PFTK1, FZD1, MTERF, AKAP9, CYP51A1, KRIT1, ANKIB1, GATAD1, ERVWE1, PEX1, and CDK6 (Supplementary Table 4). SYBR Green amplification mixture (10 ul) contained 2.5x QuantiTect SYBR Green PCR Master Mix (Qiagen), 250 nM of each forward and reverse primer, and 1 ul of cDNA as template. Reactions run on ABI PRISM 7900HT Sequence Detection System (Applied Biosystems). Cycling conditions were as follows: 10 minutes at 95°C, 40 cycles at 95°C for 15 seconds, and 60°C for 60 seconds. Glyceraldehyde phosphate dehydrogenase (GAPDH) was chosen as housekeeping gene, and commercially available primers were used (see the User Bulletin #2, Applied Biosystems, for the primers sequences). Each assay was carried out in triplicate and the transcription level was normalized using GAPDH as reference gene.

3. Results

3.1. Clinical Descriptions. The index case (Case II:2) is a 36 years old male. When he was 11 years old, he had an episode of severe headache and language disturbances. CT examination performed at that moment showed a haemorrhagic lesion in the left frontal lobe (Figure 1). Therefore, he underwent surgical treatment with good results. MRI follow-up examination was performed every two years, the last of which revealed two lesions suggestive for hemangiomas, respectively, located at the cerebellopontine angle and at the left lateral ventricle. The mother of the index case (Case I:2) is a 61 years old female. When she was 48 years old, an MRI examination of the orbit performed after an episode of right blindness associated to acute headache showed an area of haemorrhage at the superior right quadrant and at the inferior left quadrant of the right eye. She underwent surgical treatment. At present, a visual defect at the right eye still persists. The brother of the index case (Case II:1) is a 38 years old male with a history of recurrent headaches and seizures due to multiple cerebral hemorrhages from the age of 9. Surgical treatment was performed on a cerebral hemangioma located at the right parietal lobe. He underwent MRI follow-up examinations every two years. The last one revealed three lesions suggestive for hemangiomas: 1 in the right fronto parietal lobe, 1 at the cerebellopontine angle, and 1 in the cerebellum. The nephew of the index case (Case III:1) is an 11 years old male. When he was 5 years old, he reported a seizure. MRI showed two cerebral lesions suggestive for hemangiomas, one of them was bleeding. Only the father of the index case (Case I:1) showed negative results at MRI and CT examinations. In the three patients (Cases I:2, II:1, II:2) who underwent surgical treatment, the diagnosis of CCM was histologically confirmed. To date, all the family members are alive.

3.2. Genetic Analysis

3.2.1. Identification of Hemizigosity at the 7q21 Chromosome Region. Haplotype reconstruction of 13 microsatellite markers on chromosome 7q21 showed heterozygosity for markers flanking the CCM1 locus (D7S2409, D7S1813, D7S1789) in the healthy family member (I : 1), whereas all the affected members (I : 2, II : 1, II : 3, III : 1) carried and shared only the maternal haplotype, indicating a hemizygosity of the specific chromosomal region of about 700 kb (Figure 2(a)). The genetic screening of the whole coding regions of the KRIT1 gene gave negative results.

3.2.2. Identification of Complete KRIT1 and Flanking Genes Deletions by RT-QPCR. By using the RT-QPCR approach with the three primer sets described above we confirmed the complete deletion of KRIT1 gene and the hemizygous status in all affected patients (Supplementary Figure 1). Moreover, the deletion of the 5' and 3' KRIT1 flanking region was found involving the following genes: MTERF, AKAP9, CYP51A1, located downstream to the KRIT1 gene, and ANKIB1 located upstream to the KRIT1 gene (Figure 2(b)).
3.2.3. Gene Expression Analysis. To further confirm the inclusion of these genes into the genomic deletion, an expression analysis approach was set up for all genes located into the D7S2410 and D7S646 interval. FZD1 and PFTK1 were not analysed because these genes are not expressed in peripheral blood (data not shown). Results confirmed the decrease of expression of genes showing an H-CN score of 0.5 ± 0.2 and a normal expression of genes with a D-CN of 1.0 ± 0.2 (Figure 3). The ANKIB1 gene was characterized by a discrepancy between the normal DNA D-CN (detected using a couple of primers located into the intron 14) and an evident decrease of expression level (detected using a couple of primers located into the exon 6).

3.2.4. RT-QPCR Analysis for the ANKIB1 Gene Breakpoint Determination. For the ANKIB1 gene, additional RT-QPCR analysis by using intragenic primers revealed that the deletion affected a large segment of the gene with a breakpoint located in the intron 10 flanked by the amplicons ANKIB1-Ex11 (not deleted) and ANKIB1-Ex10 (deleted), (D-CN = 1.08 ± 0.03 versus H-CN = 0.53 ± 0.02, respectively, Figure 4).

4. Discussion

The use of MLPA in routine diagnostics increases dramatically the identification of large/small genomic deletion in human disease gene screening. Recently also for the three CCM associated genes, MLPA kits have been assessed and validated, and recent papers demonstrated the usefulness of this application and identified different partial or total deletions of these genes [11–15]. However, one limit of these technique, is the lack of the information about the heterozygosity of genomic region flanking the specific gene under study. This limit can be overcome by other techniques such as the RT-QPCR, or SNP Copy Number Variations
Figure 3: Copy number status Expression level analysis (± standard error mean) of genes mapping in the 7q21 chromosomal region determined by RT-QPCR was determined by RT-QPCR. Average relative mRNA expression genes level from two affected members of CCM family (II: 1 and II: 3) related to the normal ones as calibrator (1:1) to GAPDH gene as reference.

Figure 4: ANKIB1 gene breakpoint determination by RT-QPCR (± standard error mean). Results from normal genomic DNA (2 normal controls mean), from the unaffected member of family (1:1 sample), and in one affected members (II: 1) are visualized by a closed triangle (▲), a closed square (■), and a closed circle (●), respectively. The ANKIB1 exons on the left of the vertical dotted lines (exon1–10) show the copy number loss of the associated amplicons.

Assay. Although these techniques are more expensive and sometimes time consuming compared to the MLPA analysis, nevertheless, the assessment of the breakpoints allows to identify other possible deleted genes and to investigate their contribution to the disease.

Here we report a genomic deletion of about 700 kb across the CCM1 locus encompassing the MTERF, AKAP9, CYP51A1 genes, and 90% of the ANKIB1 gene. We wondered the possible relationship between the predicted function of the corresponding encoded proteins and the clinical features reported by our patients.

Few information about the function of ANKIB1 was available at the time of this work, though the ANKIB1 was recently described as a UIM (ubiquitin-interacting motif) protein with an ubiquitylation function [21]. Thus, at the moment, we are still unable to deduce if the haploinsufficiency of these proteins could influence the clinical phenotype of cavernous angiomas. The CYP51A1 gene encodes for the Sterol 14-alpha-demethylase, a member of the cytochrome P450 gene superfamily involved in sterol biosynthesis in fungi, plants, and animals [22]. There are no data about the possible involvement in human disease, and with the chemistry analysis performed on our patients.

Transcription at the 16S rRNA/tRNA (LeuUUR) gene boundary. One could ask if the stroke-like episodes and seizures, as part of symptoms showed by our patients, could overlap with those specific symptoms of MELAS diseases, rather than exclusively due to the KRIT1 gene haploinsufficiency. Unfortunately, there are no data supporting the assumption that the MELAS syndrome could be caused by the loss of expression of the MTERF protein as well as no relationship was reported between these CCM and MELAS. Thus, this hypothesis needs to be confirmed by wider epidemiological studies.

The AKAP9 gene encodes for the A-kinase anchor protein-9, a scaffolding protein that determines the subcellular localization of protein kinase A and enzymes that regulate the PKA pathway and the I(Ks) potassium channel in the heart [24]. A recent functional study described a link between genetic perturbations in AKAP9 and congenital long QT syndrome (LQTS), electrocardiographically characterized by a prolonged QT interval and polymorphic ventricular arrhythmias which may result in recurrent syncope, seizures, or sudden death [25]. However, no alteration in ECGs was observed in our CCM affected patients that could be related to this syndrome.

One notable clinical feature in our family was the anticipation of the symptoms. Although such anticipation was already described, it does not represent a common feature of the CCM and molecular mechanism remains unknown [26, 27]. It is possible that in our family the anticipation could depend on other genetic factors, mainly associated with the intrinsic genomic variability that leads the mutation
carriers to be more or less affected by haploinsufficiency of the three CCM-associated genes.

In conclusion, we report the molecular characterization of an interstitial small genomic deletion of the 7q21.2 region in a CCMs affected family, encompassing the KRIT1 gene. This paper focuses on the utility of a fine molecular characterization of this type of genomic rearrangement in CCM affected families to investigate a possible role of the flanking deleted genes. However, taking into account the absence in our survey of any other clinical features in addition to the multiple cavernous angiomas, our results can only confirm the loss of function mechanism for the already known CCM1 locus. Further evaluations of clinical features during the follow-up of our patients could contribute to clarify this issue.

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Competing Interests

The authors declare that they have no financial or personal relationships with other people or organizations that could inappropriately influence or bias this work.

References


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