The C20orf133 gene is disrupted in a patient with Kabuki syndrome

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Background: Kabuki syndrome (KS) is a rare, clinically recognisable, congenital mental retardation syndrome. The aetiology of KS remains unknown.

Methods: Four carefully selected patients with KS were screened for chromosomal imbalances using array comparative genomic hybridisation at 1 Mb resolution.

Results: In one patient, a 250 kb de novo microdeletion at 20p12.1 was detected, deleting exon 5 of C20orf133. The function of this gene is unknown. In situ hybridisation with the mouse orthologue of C20orf133 showed expression mainly in brain, but also in kidney, eye, inner ear, ganglia of the peripheral nervous system and lung.

Conclusion: The de novo nature of the deletion, the expression data and the fact that C20orf133 carries a macro domain, suggesting a role for the gene in chromatin biology, make the gene a likely candidate to cause the phenotype in this patient with KS. Both the finding of different of chromosomal rearrangements in patients with KS features and the absence of C20orf133 mutations in 19 additional patients with KS suggest that KS is genetically heterogeneous.

Abbreviations: Appr-1''-P, ADP-ribose-1''-monophosphate; Appr-1''-Pase, ADP-ribose-1''-monophosphatase; BAC, bacterial artificial chromosome; CGH, comparative genomic hybridisation; CHARGE, coloboma, heart anomalies, choanal atresia, retardation of growth and development and genital and ear abnormalities; CNV, copy-number variation; DHPLC, denaturing high-performance liquid chromatography; E, embryonic day; FISH, fluorescence in situ hybridisation; FLRT3, fibronectin-like domain-containing leucine-rich transmembrane protein 3; KS, Kabuki syndrome; NCBI, National Center for Biotechnology Information; qPCR, quantitative PCR; RT, reverse transcriptase; SMART, Simple Modular Architecture Research Tool; SNP, single nucleotide polymorphism; TEAA, triethylamine acetate; UCSC, University of California Santa Cruz
MATERIALS AND METHODS

Patients
Given the absence of validated diagnostic criteria for KS, all patients presented in this study were selected from a larger cohort and a consensus diagnosis was established by a group of experienced clinical geneticists. Informed consent was obtained from the patients or their legal representatives. Patient DNA and leukocyte suspensions were isolated following standard protocols. In total, 20 patients with KS were selected by clinical geneticists in Leuven (n = 4), Maastricht (n = 5) and Paris (n = 11).34

Patient report
The index patient with del(20)(p12.1) has been reported previously (patient 1 in Schrander-Stumpel et al). She is the second child of healthy unrelated parents and she is currently 15 years old. She was born at 39 weeks after a delivery with forceps, with a weight of 3200 g (50th–75th centile) and length of 48 cm (25th–50th centile). A left palate and severe feeding problems were present. At the age of 5 months, the left palate was surgically closed, but the feeding problems persisted. At that time, the clinical diagnosis of KS was made. Development at 1 year of age was 8 months, according to the Bayley motor and mental scale. At 14 months, her weight and height were below third centile (6.5 kg and 69.5 cm), her occipital frontal circumferences was 44.2 cm (3rd–10th centile). Development was moderately retarded. She had a premature thelarche. At 11 years, secondary sexual development started and some fat deposition around the waist was evident (weight 31.6 kg (10th–25th centile) and height 130 cm (third centile)). Cardiac ultrasound was normal. Renal investigations revealed a left sided vesico-uretral reflux grade II and an ectopic right kidney of 7.7 cm (located in the right hemipelvis, (SD 2.9)). She had habitual bladder and bowel disturbance. Ophthalmological findings included hypermetropia, bilateral astigmatism and alternating strabismus. Recurrent middle-ear infections needed transtymponal drains. She has short fifth fingers with clinodactyly and persistent fetal pads on the fingertips and pedes plani, a small urachus cyst. She is hypotonic with sialorrhoea and has striking muscle hypoplasia with relative subcutaneous fat excess, especially in the abdominal region. Her facial appearance is distinct, with long palpebral fissures and everted lower eyelids, a broad nasal tip, cleft palate, oligodontia and pre-auricular pits.

Fluorescence in situ hybridisation
FISH was performed as described previously using the P1-derived artificial chromosome clones 1043K1 and 375N15 at 8p to exclude duplications.34

Real-time quantitative PCR
Quantitative PCR (qPCR) was performed as described previously.34 Primers were designed using PrimerExpress software (Applied Biosystems, Foster City, California, USA) and analysed for the uniqueness of the sequence by using BLAT (University of California Santa Cruz (UCSC) genome browser) and BLASTN (National Center for Biotechnology Information (NCBI) database) analysis and for repeats using RepeatMasker (supplementary table 1; available online at http://jmg.bmj.com/supplemental), and PCR products were sequenced in both directions, then analysed (ABI3130; Applied Biosystems and Ensembl database, release 42).

Sequence analysis
The C20orf133 (AK131348) and the FLRT3 (a nested gene located within intron 3 of C20orf133) genes were amplified using primer sets for each exon (supplementary table 2; available online at http://jmg.bmj.com/supplemental), and PCR products were sequenced in both directions, then analysed (ABI3130; Applied Biosystems and Ensembl database, release 42).

Denaturing high-performance liquid chromatography
Denaturing high-performance liquid chromatography (DHPLC) analysis was performed according to the manufacturer’s instructions (Transgenicomics Wave system; Transgenicomics, Cheshire, UK) on genomic DNA of C20orf133 exon 14, using the primers 5’-TGA-ATC-TCA-CAT-CTC-TTA-CTA-CTT-TAT-TTC-3’ and 5’-CCA-CGC-ACA-CAC-ACA-GGT-AT-3’. A 10 μl aliquot of approximately 10 ng/μl crude PCR product was loaded onto a chromatography column (DNASEp; Transgenicomics). DNA was eluted from the column by a linear acetonitrile gradient in 0.1 mmol/l triethylamine acetate (TEAA) buffer at a constant flow rate of 1.5 ml per minute. The gradient was formed by mixing buffer A (0.1 mmol/l TEAA) and buffer B (0.1 mmol/l TEAA, 25% v/v acetonitrile). The temperature of the oven for optimal heteroduplex separation at partial DNA denaturation was deduced from melting profiles of the DNA sequence, obtained with Navigator V.1.6.2 software (Transgenicomics). Elution patterns of patient samples were compared with those of normal control samples. The total run time was 2.5 minutes and gradient conditions were 52.9–62.9°C in 2 minutes. Column temperature was 56.1°C.37

Array CGH
Array CGH at 1 Mb resolution was carried out as described previously.34 Chromosome 20 tiling path clone set was derived from type RP clones, plates 1 (offsets A1, A2, B1, B2) and 2 (offsets A1, A2, B1) and from CT type clones, plate 1 (offset A1). The plates were obtained from BACPAC Resource Center (Children’s Hospital, Oakland Research Institute, Oakland, California, USA, http://bapac.chori.org). The hybridisations and

Biosystems), in accordance with the manufacturer’s guidelines. PCR conditions were 50°C for 2 minutes, denaturation at 95°C for 10 minutes, and 40 cycles of amplification at 95°C for 15 seconds and 60°C for 1 minute. Specific PCR amplification was assessed by dissociation curve analysis. Quantification was performed using the ΔCt method and compared with the reference genes. Primers 5’-CCC-AAG-CAA-TGG-ATT-TGA-3’ and 5’-GAG-CTT-CAT-CTG-GAC-CTG-GGT-3’ in the tumour protein p53 gene, 5’-CAT-CTC-ATG-CTG-GGC-CTA-GTG-3’ and 5’-CTG-CTT-TGC-ATC-AAA-GAC-GTC-G-3’ in the annexin A3 gene (ANXA3) and 5’-TTT-TCA-TTT-TCC-TGG-CCT-TGG-3’ and 5’-TGA-CCG-CCA-GGA-GAC-AT-3’ in the olfactory receptor, family 2, subfamily B, member 2 (OR2B2) gene were used as a reference for the qPCR.

Paternity testing
Paternity testing was performed by co-amplification of the 17 autosomal STRs vWA (12p12-pter), D2S1338, TPOX (2p23.1-pter), D3S1338, FGA (4q28), D9S818, CSF1PO (5q33.3-34), D7S820, D8S1179, TH01 (11p15.5), D13S157, Penta E (15q), D16S539, D18S51, D19S433, D21S11 and Penta D (21q) as described by Decorte et al.35 36 The index patient and both parents were analysed. A statistical analysis was performed following the procedure in Decorte et al.35 36 This analysis is based on the allele frequency in a Belgian and an American population (Penta A and Penta D) and an a priori chance of 50%.
the analyses were performed as described for the 1 Mb array experiments. The threshold for an abnormal intensity ratio for a deletion is log2(3/2)–2SD. The latter is the standard deviation of all intensity ratios.

### In situ hybridisation
Mice were handled according to the guidelines of the Committee for Animal Experiments, University of Leuven, Belgium. In situ hybridisation on wild-type *Mus musculus* was performed at different stages of embryogenesis: E12.5, 14.5, 16.5 and 18.5. Embryos were isolated from pregnant mice and sacrificed by cervical dislocation. The day of plug detection was considered to be embryonic day (E)0.5. The embryos were fixed overnight in 4% paraformaldehyde in phosphate-buffered saline at 4°C and dehydrated in 70% ethanol before embedding in paraffin wax. 

Hybridisation was carried out at 70 °C for 8 hours. Post-probe. Pretreatment was performed with the mild CC2 procedure using commercial kits (Ribomap and Bluemap kits; Ventana Discovery; Ventana Medical Systems, Tucson, Arizona, USA) on an automated in situ hybridisation instrument (Ventana with digoxigenin-UTP (Roche Diagnostics, Basel, Switzerland) sections was performed with an antisense riboprobe labelled with digoxigenin-UTP (Roche Diagnostics, Basel, Switzerland) for the uniqueness of the sequence using BLAT and BLASTN analysis and for repeats using RepeatMasker software (table 1).

Quantitative reverse transcriptase PCR
Mouse adult brain and kidney, whole E11.5 embryos and mouse E18.5 tissues of liver, brain, lung, thymus, heart, intestine, bladder and kidney were used for RNA extraction (QiAquick RNA extraction kit; Qiagen, Valencia, California, USA), in accordance with the manufacturer’s protocol. A 4 μg aliquot of RNA was treated with DNase (Fermentas) before cDNA synthesis (SuperScript III; Invitrogen). cDNA was diluted four times for analysis. Primers were designed using PrimerExpress software (Applied Biosystems) and analysed for the uniqueness of the sequence using BLAT and BLASTN analysis and for repeats using RepeatMasker software (table 1).

Reverse transcriptase (RT)-qPCR was performed (qPCR MasterMix Plus for SYBR-Green I without UNG; Eurogentec, San Diego, California, USA) on an automated system (ABI7300; Applied Biosystems), in accordance with the manufacturer’s guidelines. PCR conditions were 50°C for 2 minutes, denaturation at 95°C for 10 minutes and 40 cycles of amplification at 95°C for 15 seconds and 60°C for 1 minute. Specific PCR amplification was assessed by dissociation curve analysis. Quantification was performed using the ΔCt method and compared with the reference genes. Primers 5’-GCA-ATT-CAG-CAG-GCA-ATT-AAG-3’ and 5’-ACA-ATC-GTA-CTT-AGG-GCA-GTG-ACT-3’ within the mouse homologue AK134694 gene (UCSC genome browser) and the primers 5’-ACC-CAC-GTG-CCC-ATC-TAC-3’ and 5’-AGC-CAA-GTC-CAG-AGG-AGC-3’ in the β-actin gene were designed and used as a reference for the qPCR.

### Bioinformatics
Multiple alignments of amino acid sequences was performed with ClustalW software (http://www.ebi.ac.uk/clustalw/) using the default parameter settings.

### RESULTS
FISH and 1 Mb array CGH results
In all patients, an 8p23.1 duplication, previously suggested to be causative for the KS,24 was excluded (data not shown).28 30 DNA of four patients with KS was hybridised using a bacterial artificial chromosome (BAC) array CGH at 1 Mb resolution. For three patients, the array CGH results were normal. In a fourth patient, a log intensity ratio of −0.83 of one clone (RP5-855L24, 14.55–14.69 Mb) at chromosome 20, band p12.1, was seen, indicating the presence of a deletion (data not shown). FISH analysis with RP5-855L24 on metaphase spreads of the fourth patient (hereafter referred to as the index patient) showed a signal on only one chromosome 20, thus confirming the presence of a deletion (fig 1A). FISH analysis with RP5-855L24 on metaphase spreads of the parents showed a signal on both chromosomes 20 (data not shown), suggesting a de novo deletion. Paternity testing confirmed the father to be the biological father (p<0.01).

Fine mapping the size of the deletion
To refine the size of the deletion, we analysed the DNA of the index patient using the full-tiling chromosome 20 array CGH (fig 1B). Four BACs, namely CTD-2340K11 (14.52–14.66 Mb),

### Table 1 Specific clinical features in index patient with KS, phenotypic features in KS in general and expression levels of C20orf133 in different mouse embryonic tissues and stages

<table>
<thead>
<tr>
<th>Expression of C20orf133 in mouse embryological tissue</th>
<th>Mouse (human) developmental stage</th>
<th>Phenotypic feature in index patient with KS</th>
<th>Phenotypic feature in KS</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brain: subventricular zone of striatum and olfactory lobe, cortical plate, cerebellar primordium and inferior colliculus of the tectum</td>
<td>E12.5 (6 weeks): +++; E18.5 (36 weeks): ++</td>
<td>Mental retardation, neonatal hypotonia</td>
<td>Mental retardation</td>
</tr>
<tr>
<td>Teeth: mesenchymal components of the tooth-bud condensations</td>
<td>E14.5 (7 weeks): +</td>
<td>Oligodontia</td>
<td>Hypodontia, microdontia, small dental arches</td>
</tr>
<tr>
<td>Inner ear: cells lining the vestibulocochlear and cochlear duct</td>
<td>E14.5 (7 weeks): +</td>
<td>No hearing loss</td>
<td>Hearing loss</td>
</tr>
<tr>
<td>Eye: cuboid epithelium of the lens and the inner nuclear layer of the retina</td>
<td>E16.5 (18–24 weeks): +; E18.5 (36 weeks): ++</td>
<td>Hypermetropia bilateral astigmatism, alternating strabismus, blue sclerae, ptosis</td>
<td>Colobomata and cataracts</td>
</tr>
<tr>
<td>Heart</td>
<td>E16.5 (18–24 weeks): ++; E18.5 (36 weeks): +;</td>
<td>No heart defect</td>
<td>Congenital heart disease</td>
</tr>
</tbody>
</table>

E, days post-coitum; NI, not investigated; qPCR, quantitative PCR.

+, low, but detectable, expression; ++, expression; +++, strong expression. |
C20orf133 gene disrupted in a patient with Kabuki syndrome

C20orf133 as a candidate gene for KS

The deletion in the DNA of the index patient is located within C20orf133 (AK131348 from the NITE Biological Resource Centre, or GC20P013971). This gene contains 17 exons and spans 2.06 Mb, located at 13.92–15.98 Mb. The gene structure of C20orf133 and the putative protein encoded by it is evolutionarily conserved. In particular, the N-terminal half of the putative protein is highly conserved across species. From amino acid 1–243 (of a total of 425 amino acids), the human protein shows ≥96% similarity with the Pan troglodytes, Macaca fascicularis and Mus musculus orthologues, 84% similarity with Xenopus laevis and 79% with Danio rerio. This part of the protein contains an (ADP-ribose-1''-monophosphate (Appr-1''-P) processing domain, belonging to the macro domain family. The macro domain is described in approximately 300 of the proteins expressed in the epithelium lining the lumen of the gut and stomach, and the seminiferous tubules, lungs, dorsal root ganglia and spinal cord (data not shown). In the cranial region, increased levels of C20orf133 expression were seen in the epithelial and mesenchymal components of the tooth-bud condensations, the epithelium of the primitive nasal cavity, in cells lining the vestibulocochlear and cochlear ducts, and the cranial ganglia (fig 3C and data not shown). In the E16.5 embryo, the lung and kidney maintained C20orf133 expression. Specific hybridisation was also detected in the papilla of the whisker follicles. In the eye, strong in situ staining was seen in the cuboidal epithelium of the lens and the inner nuclear (neuroblastic) layer of the retina (fig 3D and data not shown).

Mutation screening in other patients with KS

We investigated whether other patients with KS carried either deletions or point mutations in C20orf133. FISH analysis using RP5-855L24 as a probe on metaphase spreads of 14 of the 19 other patients with KS was performed and no deletion was
detected. No metaphase spreads were available for the five patients from Maastricht. Full-tiling chromosome 20 array CGH was performed on the DNA of the five patients with KS from Maastricht and the three other patients from Leuven. There was insufficient DNA from the Paris group to perform these analyses. No imbalances were detected. To exclude other mutations in the coding sequence, all C20orf133 exons and their intron–exon boundaries in the genomic DNA of the other 19 patients with KS were sequenced. A single base-pair replacement (single nucleotide polymorphism; SNP) in exon 14 (c.19C>T) was detected in 7 patients. This SNP is not present in the Human Genome Organisation SNP database. To determine whether the SNP was a common polymorphism, DHPLC was performed on 82 control DNA samples (164 alleles). The SNP was found to be heterozygous in 25 controls, and 3 controls were homozygous for the base-pair substitution. Hence, we conclude that the SNP is a benign polymorphism. Because sequencing analysis may not detect whole exon deletions or duplications that may be small enough to be below the detection limit of the current CGH array, we performed exon-specific qPCR on the available genomic DNA of the nine patients screened with the tiling array of chromosome 20. This approach confirmed the deletion of exon 5 in the index patient, but no other deletions or duplications were detected in the other nine patients.

**Mutation analysis in FLRT3**

Because genomic rearrangements and thus deletions can influence the expression of nearby genes, the effect of the detected microdeletion in the index patient might be indirect, disturbing gene function(s) in the proximity. Because neither point mutations nor deletions of the C20orf133 gene were detected in the remaining patients with KS, we investigated whether FLRT3, a nested gene located within intron 3 of the...
involved in receptor signalling protein activity, transferase activity and cell adhesion. However, neither deletions nor duplications were detected in the clones covering FLRT3 in the nine patients with KS screened using the chromosome 20 fulfilling path array. Subsequently, the three FLRT3 exons and their intron–exon boundaries were sequenced in the DNA of all 20 patients with KS, but no mutation was found.

**DISCUSSION**

We report a de novo microdeletion of \(-250\) kb at 20p12.1 in a 15-year-old girl with the KS, using array CGH with a 1 Mb resolution. This was a fortunate finding, as the resolution of the array is lower than that of the deletion size. Several observations suggest that C20orf133 is a causative gene for the observed phenotype in the index patient. First, the deletion in this patient occurred de novo in the macro-Appr-Pase-like domain of the putative protein, thereby probably disrupting the enzymatic activity. Second, the expression pattern of the C20orf133 gene during mouse embryonic development supports its importance in the development of various tissues and organs, such as those affected in this patient. The gene contains a macro functional domain, which links it with the chromatin structure. Recently, several congenital malformation syndromes have been described as caused by haploinsufficiency of a gene involved in chromatin remodelling.\(^{46-47}\)

In *Mus musculus*, both in situ hybridisation and RT-PCR showed that C20orf133 is expressed during embryonic development of the brain, in particular the ventricular zone. Just before birth, at E18.5, C20orf133 remains expressed in the brain, with relatively high levels of expression in discrete regions: the subventricular zone of the striatum and olfactory lobe, the cortical plate, the cerebellar primordium and the inferior colliculus of the tectum. The high levels of expression in the brain across a wide range of developmental stages and its persistence during adulthood is consistent with a role for the gene in mental development, and may reflect a requirement for C20orf133 in axonal outgrowth and functioning of the adult cortex. The gene is also expressed in other embryonic tissues that are typically affected in patients with KS. High levels of expression in embryonic kidney/urinary tract have been found. One study found renal malformations in 28% of tindividuals with KS,\(^{46}\) but such malformations might be underdiagnosed because they sometimes remain asymptomatic. In the current study, expression was seen in several craniofacial regions, which, given the specific facial characteristics of patients with KS, can be expected. C20orf133 was expressed in the mesenchymal components of the tooth-bud condensations at E14.5. Interestingly, dental anomalies such as hypodontia, malocclusion, microdontia and small dental arches, were seen in 68% of patients with KS in one study.\(^{46-48}\) Expression of the gene in the cells lining the vestibulocochlear and cochlear duct at stage E14.5, possibly explaining hearing loss was reported in up to 82% of patients with KS.\(^{49}\) In the developing mouse eye, strong hybridisation was seen in the cuboid epithelium of the lens and the inner nuclear (neuroblastic) layer of the retina. In situ staining on a kidney of a newborn (P0) mouse demonstrates significant expression of the mouse C20orf133 orthologue in the kidney, whereas the adrenal gland has no detectable expression.\(^{46}\)

**Figure 3** The embryonic expression pattern of the mouse orthologue of C20orf133 (AK134694). (A) Sagittal section of an E8.5 embryo within the deciduum (decid) reveals a widespread gene expression in all intra-embryonic and extra-embryonic tissues. The deciduous tissue surrounding the embryo is largely devoid of C20orf133 mRNA. (B) Transverse section (at the level of the thorax) through an E12.5 embryo showing the spinal cord and the dorsal root ganglia (DRG) shows particularly strong expression of the mouse orthologue of C20orf133 (AK134694) in the condensed DRG (red arrowhead). Lower, but still significant expression levels can be seen in the neural tube, in particular in the ventricular zone (VZ). (C) Transverse section through E14.5 kidneys shows a strong in situ signal. (D) Cranial section at E14.5. High mRNA content is seen in cells lining the vestibulocochlear and cochlear ducts, and in the cranial ganglia. (E) In the E16.5 eye, strong in situ staining can be seen in the cuboid epithelium of the lens and the inner nuclear (neuroblastic) layer of the retina. (F) In situ staining on a kidney of a newborn (P0) mouse demonstrates significant expression of the mouse C20orf133 orthologue in the kidney, whereas the adrenal gland as no detectable expression. (G) Sagittal section through an E18.5 brain reveals expression throughout the brain, with particularly high levels of expression in the subventricular zone of the striatum and olfactory lobe, the cortical plate, cerebellar primordium and the inferior colliculus of the tectum. BA, branchial arch; decid, decidium; nt, neural tube; som, somites.

C20orf133, could be affected. FLRT3 (fibronectin-like domain-containing leucine-rich transmembrane protein 3) is a transmembrane modulator of fibroblast growth factor–mitogen-activated protein kinase signalling in vertebrates and may be
post-translational modification that has a role in DNA repair, transcriptional activation and repression, telomere and chromatin biology, long-term memory formation, DNA binding, and DNA and/or RNA unwinding, among other processes. To date, no genes containing macro domains have been implicated in mental retardation and/or developmental delay. However, it is striking that several syndromes with mental retardation and multiple congenital anomalies result from haploinsufficiency of genes involved in DNA repair, transcription, chromatin biology and long-term memory formation, exactly the same processes in which macro domain-containing proteins are known to play a role. Possibly, the macro domain family of proteins may represent a novel class of dose-sensitive genes that may cause developmental disorders when mutated.

Genevieve et al reported that one of the patients with KS in their study had a clinical overlap with CHARGE syndrome. This syndrome was recently shown to be caused by mutations in or deletions of CHD7, a member of the chromodomain helicase DNA-binding genes, which have a unique combination of functional domains, including two N-terminal chromodomains, an SNF2-like ATPase/helicase domain and a DNA-binding domain. ADP-ribose binding affects the function of the Alc1 protein, a protein with homology to the Snf2 ATPase/helicase. Therefore, the alteration of the phosphorylation status of ADP-ribose by C20orf133 may have an influence on the activity of this Swi/Snf chromatin remodelling factor and thus explain the clinical overlap between some patients with CHARGE syndrome and KS.

Screening for mutations, deletions or duplications in 20 other patients with KS did not reveal any mutations within the C20orf133 candidate gene. Possibly, mutations outside the coding region as well as epigenetic changes might cause the KS phenotype. We also cannot exclude the presence of as yet unknown genes within the region, which might be affected by this deletion. However, it seems likely that KS is genetically heterogeneous, and that mutations in other genes may result in phenocopies. The accumulating findings of several different chromosomal rearrangements in patients with KS summarised above may pinpoint several loci that may cause KS.

Recently, other syndromes associated with multiple congenital anomalies and mental retardation have been shown to be genetically heterogeneous. For example, Noonan syndrome can be caused by mutations in the protein–tyrosine phosphatase non-receptor-type 11 (PTPN11) gene, or by mutations in the V-Ki Ras Sarcoma 2 (KRAS2) homologue, 46 both components of the Ras pathway, or by the V-Ki-Ras 2 Kirsten rat sarcoma 2 (KRAS2) gene.45

Noteworthy is that the haploinsufficiency of mutants in the denseosinophilic homologue 1 gene (SOS1).47 The CHARGE syndrome phenotype can, in addition to mutations in the CHD7 gene, also be caused by a mutation in the semaphoring-3E gene (SEMA3E).41 Rubinstein–Taybi syndrome can be caused by mutations in the gene encoding the protein–tyrosine phosphatase non-receptor-type 11 (PTPN11) gene, or by mutations in the Sevenless Drosophila homologue 1 gene (SOS1).47 The CHARGE syndrome phenotype can, in addition to mutations in the CHD7 gene, also be caused by a mutation in the semaphoring-3E gene (SEMA3E).41 Rubinstein–Taybi syndrome can be caused by mutations in the gene encoding the transcriptional coactivator CREBBP or by mutations in the E1A-binding protein 300-kDa (EP300) gene.

In conclusion, C20orf133 is a viable candidate gene for the phenotype in the index patient. Further evaluation is needed to determine to what extent this gene could be involved in the aetiology of KS. In addition, this study further illustrates the value of array CGH to localise genes causing developmental disorders. The fortuitous finding of a microdeletion below the 1 Mb resolution of our array demonstrates that high-resolution array CGH will enable the functional identification of more genes involved in the aetiology of KS and other clinical genetic syndromes.

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Electronic database information

NCBI accession numbers of the sequences used for alignment are NM_080676.5 (Homo sapiens), XM_001136712.1 (Pan troglodytes), AB173156 (Macaca fascicularis), CAM14292 (Mus musculus), BC060026.1 (Xenopus laevis) and NP_956843 (Danio rerio).

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