

CASE REPORT

Open Access

Complex phenotype with social communication disorder caused by mosaic supernumerary ring chromosome 19p

Caroline Demily^{1,2*}, Massimiliano Rossi^{3,4}, Gabrielle Chesnoy-Servanin^{1,2}, Brice Martin^{2,5}, Alice Poisson^{1,2}, Damien Sanlaville^{4,6} and Patrick Edery^{3,4}

Abstract

Background: Deletions or duplications of chromosome 19 are rare and there is no previous report in the literature of a ring chromosome derived from proximal 19p. Copy Number Variants (CNVs) responsible for complex phenotypes with Social Communication Disorder (SCD), may contribute to improve knowledge about the distinction between intellectual deficiency and autism spectrum disorders.

Case presentation: We report the clinical and cytogenetic characterization of a patient (male, 33 years-old, first child of healthy Portuguese non-consanguineous parents) presenting with a complex phenotype including SCD without intellectual deficiency and carrying a mosaic supernumerary ring chromosome 19p. Microarray-Based Comparative Genomic Hybridization and Fluorescence in situ Hybridization were performed. Genetic analysis showed a large mosaic interstitial duplication 19p13.12p12 of the short arm of chromosome 19, spanning 8.35 Mb. Our data suggested a putative association between psychosocial dysfunction and mosaic pure trisomy 19p13.2p12.

Conclusion: This clinical report demonstrated the need to analyze more discreet trait-based subsets of complex phenotypes to improve the ability to detect genetic effects. To address this question and the broader issue of deciphering the yet unknown genetic contributors to complex phenotype with SCD, we suggest performing systematic psychological and psychiatric assessments in patients with chromosomal abnormalities.

Keywords: Genetics, Autism, Social communication disorder, Duplication, Neurodevelopment, Chromosomal abnormalities, Trisomy, Copy number variants

Background

Deletions or duplications of chromosome 19 are rare and there is no previous report in the literature of a ring chromosome derived from proximal 19p. We report the clinical and cytogenetic characterization of a patient presenting with several abnormalities including Social Communication Disorder (SCD) without Intellectual Deficiency. This patient presented a complex phenotype with neurocognitive features, dysmorphism, growth delay and SCD. He was carrying a mosaic pure trisomy 19p13.2p12.

Pathogenic copy number variants (CNVs) are found in nearly 20% of individuals with Intellectual Deficiency (ID) [1] and in 10% of patients showing Autism Spectrum Disorders (ASD) [2]. Nevertheless, the relevance of making a clinical distinction between ID and ASD in terms of genetic etiology remains controversial because of considerable overlaps of the causative genes or chromosomal regions. Copy Number Variants (CNVs) responsible for complex phenotype with Social Communication Disorder (SCD), a diagnosis related to ASD, may contribute to improve knowledge about this data.

* Correspondence: caroline.demily@ch-le-vinatier.fr

¹Centre de dpistage et de prises en charge des troubles psychiatriques d'origine gntique, Ple Ouest, Centre Hospitalier le Vinatier, 95 bld Pinel, 69677 Bron cedex, France

²Centre de Neurosciences Cognitives, UMR 5229 (CNRS et Universit Lyon 1), Lyon, France

Full list of author information is available at the end of the article

Case presentation

Clinical report

The patient was the first child of healthy Portuguese non-consanguineous parents. The father's height was 163 cm

and the mother's was 148 cm. Family history was otherwise unremarkable. Pregnancy and delivery were normal: birth weight was 2.550 kg, at -2 standard deviations (SD); birth length was 46 cm (-3 SD); head circumference: 32 cm (-3 SD), Apgar score was 10, at 1 and 5 min. Short stature of prenatal onset was noted.

During infancy, he showed gastro-esophageal reflux and recurrent otitis media. He underwent surgical interventions for inguinal hernias, adenoidectomy and umbilical cyst ablation.

He had mild motor delay and started walking at the age of 2 years. He subsequently showed moderate learning difficulties. He has undergone several trainings in electricity and computing, but to no avail. He has had various odd jobs (e.g. waiter, fast food employee) and currently lives on his own.

He was referred to our department at the age of 33 years. Clinical data showed that he was 152 cm tall (-3.5 SD) and weighed 43.9 kg; body mass index was normal (18.9 kg/m^2) as well as head circumference (55 cm). He had a long face, high forehead, thick eyebrows, down-slanting palpebral fissures, a prominent nose with high nasal bridge and malar hypoplasia (Figure 1a and b). He had a mild scoliosis (Figure 1c) and neurological examination was normal.

He presented a psychiatric phenotype. He presented difficulties in acquiring/using language, limited effective communication and social relationships. A social

communication disorder was diagnosed according to DSM-5 criteria. The neuropsychological evaluation documented a normal intellectual functioning (total IQ: 90). The patient showed decreased psychomotor speed impacting on attentional tasks and mildly impaired verbal memory. However, he had good executive functioning and visual memory abilities.

Complete diagnostic assessment, including fragile X molecular analysis, full blood count, ammonia, plasma amino acids, urine orotic acid, screening for creatine metabolism deficiencies and urinary organic acids, was normal. Magnetic resonance imaging of the brain was normal. Skeletal survey showed mild scoliosis with no obvious sign of bone dysplasia. Echocardiography was normal and abdominal ultrasounds scan showed isolated mild hepatomegaly; liver function tests were normal.

Cytogenetic analysis

Genomic analyses were performed after obtaining a signed informed consent, according to French legislation. Those analyses are performed routinely and do not need specific ethical approval by a committee.

Chromosome analysis

Conventional blood lymphocytes karyotypes (both GTG and RHG-banding) were performed according to standard methods.

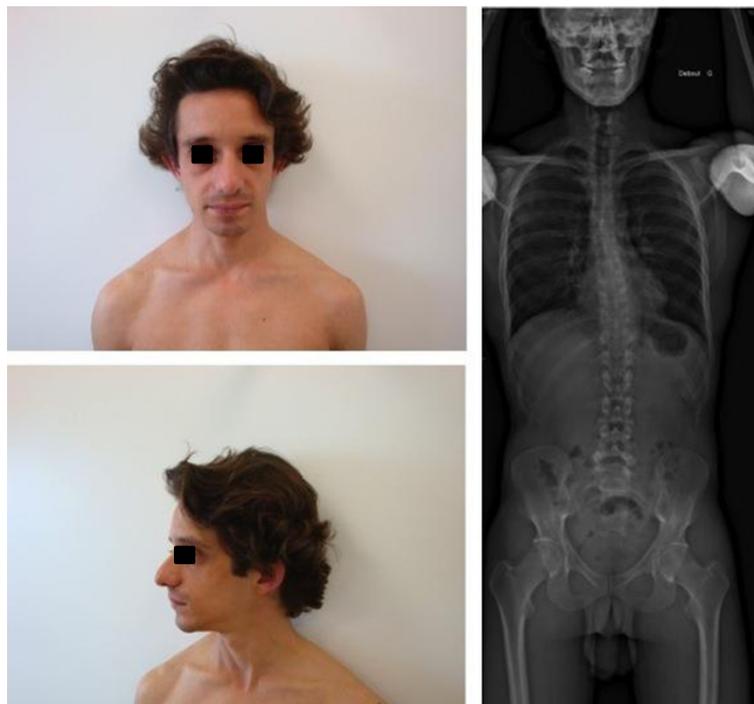


Figure 1 Facial dysmorphism of patient (a face and b profile) and X-Rays showing the scoliosis (c).

Discussion

Analysis of genomic rearrangements using aCGH in patients with various symptoms including psychiatric features exposes unexpected complexity. First of all, social communication disorder in people with chromosomal abnormalities may be much more common than reported because psychological or psychiatric assessments are not systematically carried out.

Moreover, molecular studies are usually performed for ID, ASD and/or syndromes of multiple congenital abnormalities. The genetic background of ASD is highly heterogeneous and the fact of having most common or rare CNVs may usually not be considered as a unique cause, but may occasionally increase the risk of developing ASD. Recently, Pinto et al. [5], showed that rare CNVs are an important source of risk for ASD. Also, a genome-wide screen for autism loci identified the best compatibility with linkage to 17q11.2 and 19p13, with maximum multipoint heterogeneity LOD scores of 2.9 and 2.6, respectively [6]. The mosaic gain identified with aCGH in our patient encompasses 398 genes including 97 OMIM genes and 13 morbid OMIM genes, namely *CRFL1*, *RFXANK*, *IL2RB1*, *MYO9B*, *JAK3*, *SLC5A5*, *COM*, *GDF1*, *GTPBP13*, *NDUFA13*, *INSL3*, *PIK3R2*, and *CALR3*.

To our knowledge, only 9 cases with extra ring 19chr were reported [7]. The short arm of chromosome 19 only was involved using either FISH or aCGH in two of these cases. Both patients had cerebral abnormalities including respectively enlarged cerebral ventricles and cortical atrophy [8] and Dandy Walker malformation [8]. Among these cases, only case number 19-W-p12/2-1 is suitable for comparison, because the sSMC was studied by aCGH and the genomic region included 2.53 Mb of the 19p pericentromeric region. This patient showed hip subluxation, pes calcaneovalgus congenitus, periodic breathing, congenital stridor and feeding problems. At 3 months of age, an extreme restlessness, nearly opisthotonos and at 5 months of age hyperexcitability, and developmental delay were noted. Unfortunately, social communication was not described. In Liehr's database (<http://ssmc-tl.com/sSMC.html>), 43 sSMC derived from the chromosome 19 were reported and 70% have clinical features. The polymorphic region proposed by Liehr spanned from 15.2 Mb to 39.08 Mb genomic positions, thus including the duplicated genomic region identified here. However, social cognitive aspects of individuals considered as asymptomatic with a CNV of this genomic region were not studied in detail.

Conclusion

This clinical report suggests 19p13.12p12 as a possible SCD susceptibility locus and demonstrates the need to analyze more discreet trait-based subsets of complex phenotypes to improve the ability to detect genetic effects. To address

this question and the broader issue of deciphering the yet unknown genetic contributors to complex phenotype with SCD, we suggest performing psychological and psychiatric assessments in patients with chromosomal abnormalities.

Consent

Written informed consent was obtained from the patient for publication of their individual details and accompanying images in this manuscript. The consent form is held in the patients' clinical notes and is available for review by the Editor-in-Chief.

Abbreviations

SCD: Social communication disorder; ASD: Autism spectrum disorder; CNVs: Copy number variants; ID: Intellectual deficiency; DSM-5: Diagnostic and statistical manual - Fifth revision; SD: Standard Deviation; FISH: Fluorescence In Situ Hybridization; aCGH: Microarray-based comparative genomic hybridization.

Competing interests

The authors declare that they have no competing interests.

Authors contributions

CD, MR, DS and PE designed the report. MR, AP and BM collected the clinical data. DS performed the cytogenetic analyses. GCS performed the neuropsychological assessment. CD, MS, DS and PE analysed the data and wrote the paper. All authors read and approved the final manuscript.

Acknowledgements

Nicolas Franck for giving some advices & the Conseil Scientifique de la Recherche du Vinatier.

Author details

¹Centre de diagnostic et de prises en charge des troubles psychiatriques d'origine génétique, Pôle Ouest, Centre Hospitalier le Vinatier, 95 bld Pinel, 69677 Bron cedex, France. ²Centre de Neurosciences Cognitive, UMR 5229 (CNRS et Université Lyon 1), Lyon, France. ³Hospices Civils de Lyon, service de génétique et centre de référence des anomalies du développement, GHE, Lyon, France. ⁴Centre de Recherche en Neurosciences de Lyon, Inserm U1028, UMR CNRS 5292, Université Claude Bernard Lyon 1, Lyon, France. ⁵Service Universitaire de Réhabilitation, Centre Hospitalier le Vinatier, Bron, France. ⁶Hospices Civils de Lyon, service de génétique, centre de référence des anomalies du développement, laboratoire de cytogénétique, GHE, Lyon, France.

Received: 21 July 2014 Accepted: 2 December 2014

References

1. Kaminsky EB, Kaul V, Paschall J, Church DM, Bunke B, Kunig D, Moreno-De-Luca D, Moreno-De-Luca A, Mülle JG, Warren ST, Richard G, Compton JG, Fuller AE, Gliem TJ, Huang S, Collinson MN, Beal SJ, Ackley T, Pickering DL, Golden DM, Aston E, Whitby H, Shetty S, Rossi MR, Rudd MK, South ST, Brothman AR, Sanger WG, Iyer RK, Crolla JA, et al: An evidence-based approach to establish the functional and clinical significance of CNVs in intellectual and developmental disabilities. *Genet Med* 2011, **13**:777-784.
2. Sanders SJ, Ercan-Sencicek AG, Hus V, Rui L, Murtha MT, Moreno-De-Luca D, Chu SH, Moreau MP, Gupta AR, Thomson SA, Mason CE, Bilguvar K, Celestino-Soper PBS, Choi ML, Crawford EL, Davis L, Davis Wright NR, Dhodapkar RM, DiCola M, DiLullo NM, Fernandez TV, Fielding-Singh V, Fishman DO, Frahm S, Garagaloyan R, Goh GS, Kammela S, Klei L, Lowe JK, Lund SC, et al: Multiple recurrent de novo copy number variations (CNVs) including duplications of the 7q11.23 Williams-Beuren syndrome region are strongly associated with autism. *Neuron* 2011, **70**:863-885.
3. Schluth-Bolard C, Delobel B, Sanlaville D, Boute O, Cuisset JM, Sukno S, Labalme A, Duban-Bedu B, Plessis G, Jaillard S, Dubourg C, Henry C, Lucas J, Odent S, Pasquier L, Copin H, Latour P, Cordier MP, Nadeau G, Till M, Ederly P, Andrieux J: Cryptic genomic imbalances in de novo and inherited

- apparently balanced chromosomal rearrangements: array CGH study of 47 unrelated cases. *Am J Med Genet A* 2009, **149A**:2584-2587.
4. Valli R, Marletta C, Pressato B, Montalbano G, Lo Curto F, Pasquali F, Maserati E: **Comparative genomic hybridization on microarray (a-CGH) in constitutional and acquired mosaicism may detect as low as 8% abnormal cells.** *Mol Cytogenet* 2011, **9**:4-13.
 5. Pinto D, Delaby E, Merico D, Barbosa M, Merikangas A, Klei L, Thiruvahindrapuram B, Xu X, Ziman R, Wang Z, Vorstman JA, Thompson A, Regan R, Pilorge M, Pellicchia G, Pagnamenta AT, Oliveira B, Marshall CR, Magalhaes TR, Lowe JK, Howe JL, Griswold AJ, Gilbert J, Duketis E, Dombroski BA, De Jonge MV, Cuccaro M, Crawford EL, Correia CT, Conroy J, et al: **Convergence of genes and cellular pathways dysregulated in autism spectrum disorders.** *Am J Hum Genet* 2014, **94**:677-679.
 6. McCauley JL, Li C, Jiang L, Olson LM, Crockett G, Gainer K, Folstein SE, Haines JL, Sutcliffe JS: **Genome-wide and Ordered-Subset linkage analyses provide support for autism loci on 17q and 19p with evidence of phenotypic and interlocus genetic correlates.** *BMC Med Genet* 2005, **6**:1.
 7. Melis D, Genesio R, Del Giudice E, Taurisano R, Mormile A, D'Elia F, Conti A, Imperati F, Andria G, Nitsch L: **Selective cognitive impairment and tall stature due to chromosome 19 supernumerary ring.** *Clin Dysmorphol* 2012, **21**:27-32.
 8. Novelli A, Ceccarini C, Bernardini L, Zuccarello D, Digilio MC, Mingarelli R, Dallapiccola B: **Pure trisomy 19p in an infant with an extra ring chromosome.** *Cytogenet Genome Res* 2005, **111**:182-185.

doi:10.1186/s12881-014-0132-3

Cite this article as: Demily et al.: Complex phenotype with social communication disorder caused by mosaic supernumerary ring chromosome 19p. *BMC Medical Genetics* 2014 **15**:132.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

