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TREATMENT CONTINGENT UPON ETIOLOGY:
TWO ADULT FEMALE PATIENTS WITH MILD INTELLECTUAL DISABILITY AND A CAUSATIVE COPY NUMBER VARIATION

Willem M.A. Verhoeven, Jos I.M. Egger, Nicole de Leeuw and Tjitske Kleefstra

Abstract

Objective: While modern genetic techniques have substantially increased the knowledge regarding causes of intellectual disabilities and related neuropsychiatric disorders, a marked time-lag can be noticed with respect to its implementation in common psychiatric practice. Given the relatively strong emphasis on diagnostic classification to the disadvantage of phenomenological and etiopathophysiological factors, the application of etiology-based treatment strategies may be seriously hindered. This a fortiori holds for patients with mild intellectual disability in the absence of salient dysmorphic features or congenital anomalies.

Method: The present paper illustrates this by means of two adult female patients with mild intellectual disability and late onset of symptoms from the mood and cognitive spectra.

Results and conclusions: In both patients microarray analysis disclosed de novo copy number variations, being a 2.05 Mb mosaic interstitial deletion in 7p21.1p15.3 in patient A and a 1.48 Mb interstitial duplication in 17q12 in patient B. These findings led to an appropriate diagnosis and gave direction to a significant change in treatment that resulted in an improvement of general functioning.

Key words: CNV, 7p21.1p15.3, 17q12, neuropsychiatry, cognition, etiological diagnosis

Introduction

Over the past decades, the introduction of modern genetic techniques like genomic microarray and whole exome sequencing has led to the discovery of several de novo copy number variations (CNV) and mutations providing support for their causative nature in developmental delay and intellectual disability (Vissers et al. 2010, de Ligt et al. 2012, Gilissen et al. 2014, Athanasakis et al. 2014, Martinez et al. 2017). Because of the substantially high prevalence of challenging behaviours in neuropsychiatric patients, whether or not associated with intellectual disability, the application of these techniques is of vital importance for the understanding of the disease and its etiology-guided treatment (Vulto-van Silfhout et al. 2013, McRae et al. 2017).

Since current psychiatric consultation is increasingly restricted to the inventory of symptoms as categorised in diagnostic classification systems, in this process, often, the detailed analysis of course and history including neurocognitive and social-emotional development is neglected. In this way, the importance of a psychiatric diagnosis can be easily overestimated which, in turn, may cloud important etiological factors and increase the risk of erroneous treatment decisions (Verhoeven and Egger 2014).

In this paper, two adult female patients with mild intellectual disability are described who, for the first time, developed psychiatric and cognitive symptoms around the start of their third decade.
Clinical reports

Patient A is a 39-year-old female, prematurely born from non-consanguineous parents with a birth weight of 1,380g. Apart from diabetes mellitus type II and cardiovascular problems in her mother’s family, there is no load with intellectual disability or neuropsychiatric diseases. She had one older brother who died at the age of 15 years in a serious traffic accident. When the patient was two years old she nearly drowned. Developmental milestones were all delayed and the patient tended to avoid social contact. After having followed special education, she completed domestic science school after which she was employed for unskilled working activities. Over subsequent years, the patient developed, contingent upon environmental demands, periods with pseudo-hallucinatory auditory experiences, anxieties, irritability and lowering of mood as well as inactivity and withdrawal behaviours. At the age of 23 years, she had a first contact with social psychiatry services and four years later she was hospitalized for a short period. A diagnosis of depression was made and treatment with citalopram was started, later in combination with quetiapine. Thereafter, she moved to a sheltered home facility and continued her work.

The patient was referred by her general practitioner to the specialized outpatient department for neuropsychiatry for re-evaluation of mood complaints in the context of her delayed development.

At examination, a female obese patient was seen with large ears and lobules, deep set eyes and columella below alae nasi, but no major dysmorphisms. Height, weight and head circumference were 174cm (+0.5 SD), 124kg (>-2 SD) and 55.5cm (0 SD), respectively. Neuropsychiatric and somatic/neurological examination disclosed a tendency to referential thinking, but no signs of major depression or other abnormal findings. Haematological and biochemical parameters including FT4 were all within normal range. Extensive neuropsychological testing revealed mild intellectual disability (WAIS-IV Total IQ: 54) with lowered verbal, nonverbal and social memory, and lowered speed of information processing and executive functioning. Cytogenetic analysis revealed no changes as compared to six years before but no signs of distress, psychiatric symptoms or psychotherapeutic interventions were started.

The patient was referred again because of suspected gradual decline of neurocognitive functioning as well as complaints about increased anxiety levels. Intellectual capacities were similar to the former measurement (WAIS-IV Total, Verbal and Performal IQ: 53, 51 and 72, respectively). As to neurocognition, performance levels were generally comparable to those found ten years before except for speed of information processing and executive functioning which were both lowered. In addition, some indication for socio-emotional overestimation could not be ascertained. MRI scanning of the brain revealed no changes as compared to six years before and was considered normal. Genome wide CytoScan HD array analysis showed a paternally inherited 205 kb gain in 22q11.1 and a de novo, interstitial duplication of 1.48 Mb in 17q12 encompassing 24 genes (arr[hg19]17q12(34,823,616-36,307,774)x3 pat; Figure 2). The duplication in 17q12 is a known recurrent microduplication that is associated with the so-called 17q12 duplication syndrome (OMIM #614526) which is predominantly characterized by mild to moderate intellectual disability.

All reported complaints about cognitive functioning and recurrent anxieties could be emanated from enhanced susceptibility to context-related socio-emotional overestimation. Therefore, a more structured environmental approach was recommended with periodic re-evaluation of cognitive functioning. No further psychopharmacological or psychosocial interventions were indicated.

Discussion

In this paper, two adult female patients with mild...
Figure 1. A. Array plot of chromosome 7 of patient A with a de novo, mosaic, interstitial deletion of 2.05 Mb in 7p21.1p15.3. The single horizontal line represents the probe signal values, with n=2 for a normal copy number. The genotype information is represented below this single line by the so-called B-Allele Frequency, with three combinations possible in case of n=2 (AA, BA or BB) and two combinations for n=1 (A or B), if there is a deletion. In case of a mosaic loss there will be a mixture of cells with and without loss of the 7p21.1p15.3 allele resulting in four possible combinations (BB, BA, AB or AA). The plot in blue shows the array findings in DNA from buccal swab cells and the plot in pink the array findings in DNA from peripheral blood. B. Screen shot of the UCSC genome browser (https://genome.ucsc.edu) of the aberrant 7p21.1p15.3 region encompassing 12 protein-coding genes. The probe coverage on the CytoScanHD array platform and the presence of repeat sequences are indicated in the lower part of the figure.
Figure 2. A. Array plot of chromosome 17 of patient B with a de novo, interstitial gain of 1.48 Mb in 17q12. The single horizontal line represents the probe signal values, with n=2 for a normal copy number. The genotype information is represented below this single line by the so-called B-Allele Frequency, with three combinations possible in case of n=2 (AA, BA or BB), two combinations for n=1 (A or B), if there is a deletion, and four combinations in case of a gain, n=3 (AAA, AAB, ABB or BBB).

B. Screen shot of the UCSC genome browser (https://genome.ucsc.edu) of the aberrant 17q12 region encompassing 24 protein-coding genes. The probe coverage on the CytoScanHD array platform and the presence of repeat sequences are indicated in the lower part of the figure.
intellectual disability and relatively adequate social participation are described. In both, de novo CNVs were demonstrated to be causatively implicated in their developmental delay. In patient A, a mosaic interstitial deletion in 7p21.1:p15.3 was detected while in patient B a 17q12 microduplicaton was established. In both patients, referral was initiated because of suspected cognitive decline and/or psychiatric symptoms from the affect domain.

With respect to the 7p deletion, no deletions of comparable size have been reported so far. There are only a few patients described in the literature and a mere dozen patients registered in clinical genetic databases such as ECARUCA (www.ecaruca.net) and DECIPHER (https://decipher.sanger.ac.uk). However, these concern patients with significantly larger deletions that are associated with severe intellectual disability, marked dysmorphisms and somatic anomalies (Chotai et al. 1994, Shimada et al. 2013).

The 17q12 duplication syndrome is associated with a highly variable degree of intellectual disability and, in most cases, with neurological and somatic disorders, such as seizures and renal abnormalities (Mefford et al. 2007, Nagamani et al. 2010, Caselli et al. 2010, Faguer et al. 2011, Bierhals et al. 2013).

As to patient A, more than 15 years prior to referral a diagnosis of major depression was made after which she underwent a form of psychotherapy focused on anxious cognitions. In addition, she was treated over a long period of time with psychotropics. The psychotherapeutic intervention did not attenuate her anxious feelings and affective dysregulations and was therefore discontinued. Psychopharmacological treatment lacked any effect and resulted in morbid obesity (BMI: 41). Recurrent complaints about cognitive functioning brought patient B under the attention of a neurologist who did not detect any abnormalities and forwarded her to a psychotherapist to cope with anxieties and feelings of bereavement related to her mother’s death. No improvement was achieved, instead, symptoms of distress and memory complaints intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified. In both patients, personalized contextual measures with increased staff involvement resulted in an intensified.

In conclusion, as can be inferred from the above, the absence of notable facial dysmorphisms or somatic abnormalities often does not signal the clinician to the absence of notable facial dysmorphisms or somatic abnormalities often does not signal the clinician to the absence of notable facial dysmorphisms or somatic abnormalities often does not signal the clinician to the absence of notable facial dysmorphisms or somatic abnormalities often does not signal the clinician to the absence of notable facial dysmorphisms or somatic abnormalities often does not signal the clinician to the absence of notable facial dysmorphisms or somatic abnormalities often does not signal the clinician to the absence of notable facial dysmorphisms or somatic abnormalities often does not signal the clinician to.

Acknowledgments

Written informed consent for publication of histories and genetic findings was obtained from patients A and B and their parents. A copy of the written informed consent is available for review by the Editor-in-Chief of this journal. The authors are indebted to the patients and their parents for their kind cooperation. The authors declare that they have no competing interests. All authors read and approved the final manuscript.

Patient A and B are included in the ECARUCA database with ID 5204 and 5225, respectively. This study is part of a collaborative project of the research group ‘Psychopathology and Genetics’ of the Radboud University Nijmegen and the Vincent van Gogh Institute for Psychiatry, Venray, the Netherlands.

References


