

Cognitive, Behavioral, and Adaptive Profiles in Williams Syndrome With and Without Loss of *GTF2IRD2*



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Carlos Alberto Serrano-Juárez,¹ Carlos Alberto Venegas-Vega,² Ma. Guillermina Yáñez-Téllez,¹ Mario Rodríguez-Camacho,¹ Juan Silva-Pereyra,¹ Hermelinda Salgado-Ceballos,³ AND Belén Prieto-Corona¹

¹Laboratorio de Neurometría, Facultad de Estudios Superiores Iztacala, UNAM, Los Reyes Iztacala, Tlalnepantla, Estado de México, CP

²Servicio de Genética, Hospital General de México “Dr. Eduardo Liceaga”, Cuauhtémoc, CDMX, CP

³Unidad de Investigación Médica en Enfermedades Neurológicas, Hospital de Especialidades, CMN “Siglo XXI”, IMSS

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Abstract

Williams syndrome (WS) is a neurodevelopmental disorder that results from a heterozygous microdeletion on chromosome 7q11.23. Most of the time, the affected region contains ~1.5 Mb of sequence encoding approximately 24 genes. Some 5–8% of patients with WS have a deletion exceeding 1.8 Mb, thereby affecting two additional genes, including *GTF2IRD2*. Currently, there is no consensus regarding the implications of *GTF2IRD2* loss for the neuropsychological phenotype of WS patients. **Objectives:** The present study aimed to identify the role of *GTF2IRD2* in the cognitive, behavioral, and adaptive profile of WS patients. **Methods:** Twelve patients diagnosed with WS participated, four with *GTF2IRD2* deletion (atypical WS group), and eight without this deletion (typical WS group). The age range of both groups was 7–18 years old. Each patient’s 7q11.23 deletion scope was determined by chromosomal microarray analysis. Cognitive, behavioral, and adaptive abilities were assessed with a battery of neuropsychological tests. **Results:** Compared with the typical WS group, the atypical WS patients with *GTF2IRD2* deletion had more impaired visuospatial abilities and more significant behavioral problems, mainly related to the construct of social cognition. **Conclusions:** These findings provide new evidence regarding the influence of the *GTF2IRD2* gene on the severity of behavioral symptoms of WS related to social cognition and certain visuospatial abilities. (*JINS*, 2018, 24, 896–904)

Keywords: Williams syndrome, Visuospatial abilities, Neuropsychological phenotype, Neuropsychology, Social cognition, Chromosomal microarray

INTRODUCTION

Williams syndrome (WS) is a rare neurodevelopmental disorder caused by a microdeletion in the q11.23 region of chromosome 7. The incidence rate of its typical forms is 1/7000 to 1/25,000 live births (Bellugi, Lichtenberger, Mills, Galaburda, & Korenberg, 1999; Mervis & Morris, 2007; Meyer-Lindenberg, Mervis, & Berman, 2006). The main clinical manifestations of WS are: (1) characteristic facial features (e.g., small upturned nose, long philtrum, wide mouth, and small chin); (2) supravalvular aortic stenosis (SVAS) and/or peripheral pulmonary artery stenosis (PPAS); and (3) an atypical neuropsychological phenotype characterized by a moderate to

severe intellectual disability, severe impairment of visuospatial function, mild language impairment (Ferrero et al., 2010), and hypersociability (Bellugi et al., 1999). Most commonly (91–95%), WS is a result of deletion of 1.5 megabases (Mb) spanning a total of 24 genes. Near the middle of this deletion span is *ELN*, the gene that encodes elastin, mutations of which are associated with the SVAS/PPAS aspect of the syndrome (Mervis & Morris, 2007; Osborne, 2010; Porter et al., 2012).

The WS neuropsychological profile is strongly associated with a severe impairment in visuospatial capacity (Capirci, Sabbadini, & Volterra, 1996; Garayzábal, 2005; Garayzábal & Cuetos, 2008). This visuospatial impairment could be consequent to the loss of genes involved in the structure and functions of the parietal cortex and the so-called dorsal stream of processing thought to carry location and motion information anterolaterally from primary visual cortex to parietal areas (Atkinson et al., 2001; Atkinson & Braddick, 2012;

Correspondence and reprint requests to: Belén Prieto-Corona, Av. De los Barrios No. 1, Los Reyes Iztacala, Tlalnepantla, Edo. Méx. 54090, México. E-mail: bemapado@gmail.com

Atkinson & Nardini, 2008; García-Nonell, Rigau-Ratera, Artigas-Pallarés, García-Sánchez, & Estévez-González, 2003). Some authors have found gray matter volume alterations or reductions in the dorsal occipitoparietal sulcus/vertical region of the intraparietal sulcus, which could explain, at least in part, dorsal stream vulnerability in WS (Atkinson, 2017; Atkinson & Braddick, 2011; Meyer-Linderberg et al., 2004).

There has been increasing recognition of the variability in WS cognitive phenotype, including variability in intelligence (Tassabehji et al., 1999), executive functions (Morris et al., 2003), the expression of autistic-like features (Edelmann et al., 2007), and visuospatial abilities (Karmiloff-Smith et al., 2003). Advancements in genetic and molecular biology techniques, such as chromosomal microarrays (CMAs), can enable genomic alterations to be identified with higher precision than previously. Hence, the cognitive phenotype can be analyzed in combination with CMA data to determine the genes affected by each patient's deletion and thus help to explain the variability observed in the neuropsychological phenotypes of WS patients.

Studies conducted in animal models, as well as in patients with atypical deletions, have identified genes within the range of the WS microdeletion that could affect the neuropsychological phenotype of WS, including, from centromere to telomere (see Figure 1), *FZD9*, *BAZ1B*, *STX1A*, *LIMK1*, *CLIP2*, *GTF2IRD1*, and *GTF2I* (Botta et al., 1999; Frangiskakis et al., 1996; Gray, Karmiloff-Smith, Funnell, & Tassabehji, 2006; Morris et al., 2003; Osborne, 2010; Vandeweyer, Van der Aa, Reyniers, & Kooy, 2012; Wang, Spörle, Paperna, Schughart, & Francke, 1999). These genes are known regulators of proteins related to localized structural and functional brain development (Broadbent et al., 2014; Gao et al., 2010; Karmiloff-Smith et al., 2012, 2003; Osborne, 2010; Tassabehji et al., 1999; Vandeweyer et al., 2012).

Most cases of WS (91–95%) are associated with loss of *GTF2I* and *GTF2IRD1*, which are normally expressed in the cerebellum, hippocampus (Chailangkarn, Noree, & Muotri, 2018), and intraparietal sulcus (Hoeft et al., 2014). Disruption of

these genes has been associated with reduced fear and aggression (Young et al., 2008), altered visuomotor integration and visuospatial processing (Hirota et al., 2003; Hoeft et al., 2014), intellectual disability (Morris et al., 2003), and highly sociable behavior (Chailangkarn et al., 2018; Young et al., 2008).

A small minority (5–8%) of patients with WS present with an atypical 1.8-Mb deletion that encompasses two additional genes, including *GTF2IRD2*, a member of the *GTF2I* gene family, whose members encode transcription factors required for synapse development and neurulation (Allen Brain Atlas, 2010; Makeyev et al., 2004; Tipney et al., 2004; Uhlén et al., 2015). The *GTF2I* gene family also includes *BEN*, a *TFII-I* family gene (Li et al., 2015). In mice, haploinsufficiency of *TFII-I* proteins results in a variety of phenotypic manifestations, including embryonic lethality and brain hemorrhage, as well as vasculogenic, craniofacial, and neural tube defects (Enkhmandakh et al., 2009).

GTF2IRD2 mRNA expression levels are high in the cerebral cortex, specifically in the prefrontal and parietal areas, as well as in the cerebellum (Allen Brain Atlas, 2010; Porter et al., 2012; Tipney et al., 2004; Uhlén et al., 2015). The protein encoded by *GTF2IRD2* is also a *TFII-I* family protein (Makeyev et al., 2004) expressed in the brain (Tipney et al., 2004), specifically in parietal, frontal (orbitofrontal and dorsolateral) cortices and in the cerebellum (Allen Brain Atlas, 2010; Porter et al., 2012; Uhlén et al., 2015). *GTF2IRD2* has been shown to regulate the activity of *GTF2IRD1* and other *TFII-I* proteins during the postnatal period by direct interaction and sequestration of the proteins in a nuclear compartment (Palmer et al., 2012).

The aforementioned evidence indicates that disruption of *GTF2I* family genes in WS is associated with altered social behavior, negative emotionality, visuospatial ability, and intellectual ability (Crespi & Hurd, 2014; Hoeft et al., 2014; Morris et al., 2003; Young et al., 2008). Meanwhile, *GTF2IRD2*, specifically, has been associated with visuospatial functioning, social reasoning, and cognitive flexibility (Porter et al., 2012). Therefore, we hypothesized that loss of

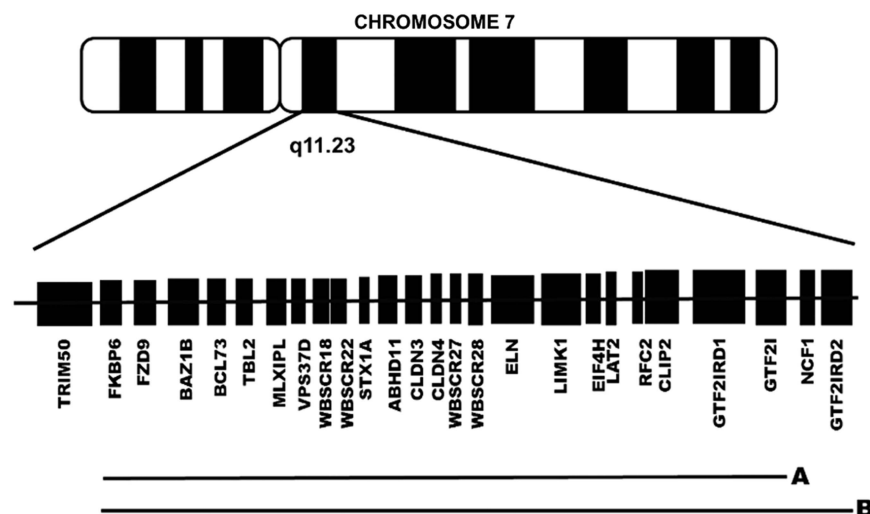


Fig 1. Schematic representation of the genes lost in the typical group of patients with a preserved *GTF2IRD2* (1.5 Mb deletion) (A) and in the atypical group of patients without *GTF2IRD2* (1.8 Mb deletion) (B).

GTF2IRD2 may underlie visuospatial and social skill impairment severity in WS patients. To test this hypothesis in this study, we evaluated the cognitive, behavioral, and adaptive effects of *GTF2IRD2* deletion in WS patients.

METHODS

Participants

Twelve patients (7 males) diagnosed with WS (age range, 7–18 years) were recruited from the *Asociación Nacional de Síndrome de Williams A.C.*, the *Asociación Viviendo con Síndrome de Williams A.C.*, and from the Genetics Service of the “Dr. Eduardo Liceaga” General Hospital of Mexico. All patients were living in the metropolitan area of Mexico City. Before starting the assessments, the parents of the participants signed written informed consent forms, and the patients’ participation was approved by the ethics committee of the Faculty of Higher Education Iztacala, which adheres to the Helsinki Declaration.

Based on the CMA results (described below), patients were divided into a *typical GTF2IRD2*-retained group and an *atypical GTF2IRD2*-deleted group (Figure 1). The typical group consisted of 8 patients (5 males) with a 1.5-Mb microdeletion and a chronological age (CA) mean of 11.5 years (*SD*, 3.59), including five who were in a special education program and four who had a perinatal complication. The atypical group consisted of 4 patients (2 males) with a 1.8-Mb microdeletion and a mean CA of 12 years (*SD*, 4.97), including one who was in a special education program, one who had a perinatal complication. None of the patients in either the typical group or the atypical group were taking medication. As shown in Supplementary Table 1S, the two groups were similar with respect to CA, mental age, full-scale IQ (FSIQ), and number of completed years of education (Mann Whitney *U*, all $p > .05$). Exact Fisher tests revealed no significant between-group differences with respect to gender constitution ($p = .57$), special education rate ($p = .27$), and perinatal complication rate ($p = .42$).

Setting and Procedures

A geneticist conducted a clinical assessment of each participant and evaluated his or her CMA results. A trained neuropsychologist conducted a neuropsychological evaluation in two 1.5-hr sessions in a cubicle at the Genetics Services of the “Dr. Eduardo Liceaga” General Hospital of Mexico. The CMA results were revealed to the neuropsychologist after all neuropsychological tests were completed. All assessments were conducted in Spanish, and Spanish-language neuropsychological instruments were used.

Genetic Assessment

In the geneticist’s office, a blood sample was obtained from each participant for CMA Cyto-Scan Optima analysis to

determine his or her 7q11.23 microdeletion scope. This genetic analysis technique is based on nucleic acid hybridization and fluorescence analysis of imaged chromosomes (Venegas Vega, 2012).

Neuropsychological Instruments

To obtain each participant’s clinical history, a structured interview was carried out to collect information about each patient’s personal (pathological and non-pathological), hereditary-familial, and developmental history. The Wechsler Intelligence Scale for Children, 4th edition (WISC-IV; Wechsler, 2007) and the Wechsler Adult Intelligence Scale, 4th edition (WAIS-IV; Wechsler, 2014) were administered to evaluate the intelligence of patients who were ≤ 17 years of age and of patients who were 17–18 years of age, respectively. The following WISC-IV/WAIS-IV scales were considered: Verbal Comprehension Index (VCI), Perceptual Reasoning Index (PRI), Working Memory Index (WMI) and Processing Speed Index (PSI), and FSIQ. Attention, memory, and executive function were evaluated with the NEUROPSI Attention and Memory subtests (Ostrosky-Solís, Guevara-López, & Matute, 2012) and the Child Neuropsychological Assessment 2 (Evaluación Neuropsicológica Infantil, ENI-2) (Matute, Rosselli, & Ardilla, 2014). Visual-spatial processing ability was assessed with the Developmental Test of Visual Perception, 3rd edition (DTVP3), which uses the Frostig perception evaluation method (Hammill, Pearson, & Voress, 2016).

To detect a wide range of emotional and behavioral problems, parents of the WS patients completed the family questionnaire of the Assessment System for Children and Adolescents (Sistema de Evaluación de Niños y Adolescentes (SENA) (Fernández-Pinto, Santamaría, Sánchez-Sánchez, Carrasco, & Del Barrio, 2015). The SENA measures internalizing problems (e.g., depression, anxiety, social anxiety), externalizing problems (e.g., hyperactivity and impulsivity, attention problems, aggressiveness), specific problems (e.g., developmental delays, learning disabilities), areas of vulnerability, and psychological resources. Finally, parents also completed the Adaptive Behavior Assessment System, 2nd edition (ABAS-II) parent form (Harrison & Oakland, 2008); the ABAS-II provides a complete assessment of adaptive skills across the lifespan.

Statistical Analysis

Mann Whitney *U* tests were used to evaluate intergroup differences in neuropsychological test scores because the cohort was too small to produce parametric datasets. Due to this study’s the small size, no *p*-level adjustment for multiple comparisons was made. Nevertheless, we calculated Cohen’s *d* statistical values for each between-group comparison result, such that $d = 0.2$ was considered to represent a small effect, 0.5 a medium effect, and 0.8 a large effect. We analyzed the

Table 1. Comparison of DTVP-3 between atypical and typical WS group

Variable	Typical group			Atypical group			Mann Whitney U		Cohen's <i>d</i>
	<i>M</i>	<i>SD</i>	Mean rank	<i>M</i>	<i>SD</i>	Mean rank	U	<i>p</i> -Value	
Visual-motor integration index	59.17	13.39	5.67	57.50	14.15	5.25	11.00	.91	0.14
Eye-hand coordination	2.17	1.94	5.83	2.25	2.50	5.00	10.00	.76	0.27
Copying	4.33	2.80	5.83	3.75	2.63	5.00	10.00	.76	0.27
Reduced motor response index	85.00	4.34	7.08	73.75	9.95	3.13	2.50	.04	1.67 ^a
Figure-ground	8.17	1.83	5.83	7.75	2.22	5.00	10.00	.76	0.27
Visual closure	3.50	0.84	7.17	2.00	0.82	3.00	2.00	.04	1.83 ^a
Form constancy	10.83	1.17	7.00	7.50	3.32	3.25	3.00	.07	1.53 ^a
General visual perception index	74.17	7.22	6.33	67.00	9.83	4.25	7.00	.35	0.72

Note: Two participants did not perform the DTVP-3 because they did not return.

^aLarge effect size.

standardized scores obtained on all administered neuropsychological instruments.

RESULTS

Participants in both groups obtained FSIQ scores in the range of 40 to 60. No intergroup differences were found in any of the WISC-IV/WAIS-IV indices, nor in the NEUROPSI attention and memory test or the ENI-2 Executive Function scale. The two groups had similar verbal ability scores. On the other hand, a tendency toward lower cognitive performance was observed in the atypical *GTF2IRD2*-deleted group, relative to the typical group, especially in the tests that assess cognitive flexibility. The results of the intergroup comparison can be seen in Supplementary Tables S2–S4.

In DTVP-3 (Table 1), the typical group ($\bar{X} = 3.5$; $SD = 0.84$) performed better on the Visual Closure subtest than the atypical group ($\bar{X} = 2.00$; $SD = 0.82$), with a large effect size ($U = 2$; $p = .04$; $d = 1.83$). The typical group ($\bar{X} = 10.83$; $SD = 1.17$) showed a non-significant trend toward performing better than the atypical group ($\bar{X} = 7.5$; $SD = 3.32$) on the Form Consistency subtest, with a large effect size ($U = 3$; $p = .07$; $d = 1.53$). Relative to the typical group ($\bar{X} = 85$; $SD = 4.34$), the atypical group ($\bar{X} = 73.75$; $SD = 9.95$) had greater alterations in the Reduced Motor Response Index, of which the Visual Closure and Form Consistency subtests are components, with a large effect size ($U = 2.5$; $p = .04$; $d = 1.67$). Note that for this index perceptual abilities are highly weighted given the limited use of motor skills in the execution of its subtests.

In the SENA inventory, the atypical group ($\bar{X} = 80.75$; $SD = 10.72$) scored higher than the typical group ($\bar{X} = 59.75$; $SD = 15.58$) on the Isolation scale, with a large effect size ($U = 4$; $p = .04$; $d = 1.46$), suggesting that atypical patients spend more time alone (Table 2). In the same way, the typical group ($\bar{X} = 52.88$; $SD = 9.51$) performed better than the atypical group ($\bar{X} = 36.25$; $SD = 3.40$) on the Emotional Intelligence scale of the SENA, with a large effect size ($U = 1$; $p = .01$; $d = 2.17$), evidencing better social skills and behaviors of the typical group compared to the

atypical group. Analyzing the items that make up the SENA Emotional Intelligence and Isolation scales, we found that, compared to the typical group, the atypical group present greater deficiencies in several constructs of social cognition (Adolphs, 2009), including emotional recognition (e.g., “When I feel bad, he/she notices”), empathy (e.g., “Pay attention to how others feel”), and theory of mind (e.g., “Knows how to put themselves in the another’s place”).

Finally, the typical group ($\bar{X} = 6.75$; $SD = 3.06$) also performed significantly better than the atypical group ($\bar{X} = 2.25$; $SD = 2.50$) on the ABAS-II Leisure scale, with a large effect size ($U = 4$; $p = .04$; $d = 1.46$). Furthermore, the typical group ($\bar{X} = 8.88$; $SD = 2.10$) also performed better on the ABAS-II Social scale than the atypical group ($\bar{X} = 5$; $SD = 2.71$), with a large effect size ($U = 3$; $p = .02$; $d = 1.65$). The poorer social skills of the atypical group relative to the typical group are consistent with a greater deficit in social cognition, which could be related to the loss of *GTF2IRD2* (Table 3).

DISCUSSION

Our typical group with loss of *GTF2I/GTF2IRD1* showed a neuropsychological phenotype characteristic of WS, whereas our atypical group, whose 1.8-Mb deletion resulted in the additional loss of *GTF2IRD2*, exhibited certain visuospatial/perception and social cognition alterations together with a reduced ability to socialize. *GTF2IRD2* has been observed to have a genetic regulatory influence on *GTF2I* and *GTF2IRD1* (Palmer et al., 2012); all three of these genes are expressed in frontal and parietal cortices, and the cerebellum (Allen Brain Atlas, 2010; Porter et al., 2012; Tipney et al., 2004; Uhlén et al., 2015). Considering the present results, we posit that the loss of this group of genes, and the consequent loss of the availability of the proteins encoded by these genes in the prefrontal cortex, parietal cortex, and cerebellum, where they are normally highly expressed, led to greater cognitive impairments than is typical of WS.

We found that both groups had similar cognitive-behavioral phenotypes with respect to verbal abilities. This

Table 2. Comparison of SENA between atypical and typical WS group

Variable	Typical group			Atypical group			Mann Whitney U		
	<i>M</i>	<i>SD</i>	Mean rank	<i>M</i>	<i>SD</i>	Mean rank	U	<i>p</i> -Value	Cohen's <i>d</i>
Index scores									
Global problems index	54.50	16.48	6.25	53.00	4.76	7.00	14.00	.81	0.27
Emotional problems index	52.13	15.67	6.25	49.25	4.99	7.00	14.00	.81	0.27
Behavioral problems index	54.13	14.33	6.38	52.75	5.56	6.75	15.00	.93	0.41
Executive functions index	55.75	11.99	6.00	57.25	4.03	7.50	12.00	.57	0.40
Personal sources index	44.38	11.56	7.38	36.25	8.02	4.75	9.00	.28	0.73
Clinical scales									
Depression	57.13	22.74	6.56	51.75	7.54	6.38	15.50	.93	0.05
Anxiety	57.25	16.26	6.50	52.50	3.32	6.50	16.00	1	0.00
Social anxiety	42.00	9.56	5.63	45.50	6.14	8.25	9.00	.28	0.73
Somatic complaints	50.50	7.86	6.56	48.25	8.96	6.38	15.50	.93	0.05
Attention problems	58.38	9.62	6.31	61.50	4.80	6.88	14.50	.81	0.15
Hyperactivity	53.75	8.55	5.94	55.75	3.10	7.63	11.50	.46	0.45
Anger management problems	56.50	17.05	6.94	49.50	10.38	5.63	12.50	.57	0.35
Aggression	52.25	12.97	5.88	54.50	9.81	7.75	11.00	.46	0.51
Behavioral disruptive	50.75	9.78	6.25	53.75	1.26	7.00	14.00	.81	0.20
Unusual behavior	63.25	14.85	6.25	69.00	24.15	7.00	14.00	.81	0.20
Vulnerability scales									
Emotional regulation problems	56.25	13.99	6.63	53.50	7.42	6.25	15.00	.93	0.10
Rigidity	49.13	9.93	6.00	52.00	7.35	7.50	12.00	.57	0.40
Isolation	59.75	15.58	5.00	80.75	10.72	9.50	4.00	.04	1.46 ^a
Personal sources scales									
Social integration	45.75	10.51	7.31	38.75	9.64	4.88	9.50	.28	0.67
Emotional Intelligence	52.88	9.51	8.38	36.25	3.40	2.75	1.00	.01	2.17 ^a
Disposition to the study	38.75	17.43	6.44	40.50	5.80	6.63	15.50	.93	0.05

^aLarge effect size.**Table 3.** Comparison of ABAS-II between atypical and typical WS group

Variable	Typical group			Atypical group			Mann Whitney U		
	<i>M</i>	<i>SD</i>	Mean rank	<i>M</i>	<i>SD</i>	Mean rank	U	<i>p</i> -Value	Cohen's <i>d</i>
Areas									
Communication	3.88	3.23	6.69	2.75	1.71	6.13	14.50	.81	0.15
Academic skill	1.00	0.00	6.00	1.50	1.00	7.50	12.00	.57	0.40
Self-direction	6.50	3.66	7.69	3.00	2.45	4.13	6.50	.11	1.05
Leisure	6.75	3.06	8.00	2.25	2.50	3.50	4.00	.04	1.46 ^a
Social	8.88	2.10	8.13	5.00	2.71	3.25	3.00	.02	1.65 ^a
Use of community resources	3.88	3.68	6.63	3.50	3.00	6.25	15.00	.93	0.10
Home life	7.50	3.16	6.94	6.50	3.87	5.63	12.50	.57	0.35
Health and security	6.00	3.59	7.44	3.25	2.63	4.63	8.50	.21	0.79
Self-care	4.50	2.78	7.50	2.50	1.29	4.50	8.00	.21	0.85
Index scores									
Conceptual index	66.63	9.61	7.38	59.50	6.56	4.75	9.00	.28	0.73
Social index	86.88	9.75	8.13	67.25	12.69	3.25	3.00	.02	1.65 ^a
Practical index	72.38	13.36	7.44	62.25	8.22	4.63	8.50	.21	0.79
General adaptive coefficient	71.63	11.21	7.69	61.50	8.27	4.13	6.50	.10	1.05

^aLarge effect size.

sparing of verbal abilities may be related to the low-density of expression of *GTF2IRD2* in temporal areas (Allen Brain Atlas, 2010; Colantuoni et al., 2011; Uhlén et al., 2015;

Zhang et al., 2014). Meanwhile, we found that *GTF2IRD2* deletion in WS was associated with impairments in visuospatial skills and social cognition. These results suggest that

GTF2IRD2 may contribute to the severity of the neuropsychological phenotype of WS, especially with respect to deficiencies in some visuospatial abilities and some aspects of social cognition.

Concerning visuospatial ability, patients with a *GTF2IRD2* deletion showed reduced visual closure capacity, compared to patients in whom the gene was conserved. This ability is related to recognition of how components of a unit are associated to form a figure (Frostig, 1999). Formerly, Porter et al. (2012) found low scores on spatial tasks in nine WS patients who lost this gene. However, in the present study, we found specifically that visual closure ability, form consistency, and figure-grounding, tasks that do not include a motor component and that form the reduced motor response index, are weaknesses in these patients. These findings support the notion that the *GTF2IRD2* deletion phenotype may have more of a perceptual component than a visual-motor one.

Additionally, visual closure capacity has been found to be related to global attention tasks (D'Souza, Booth, Connolly, Happe, & Karmiloff-Smith, 2016; Porter & Coltheart, 2006). Dorsal stream vulnerability theory encompasses visual closure capacity together with navigation capacity, use of spatial labels, and visuoconstructive (Atkinson, 2017; Atkinson et al., 2001; Atkinson & Braddick, 2011, 2012; Atkinson & Nardini, 2008). According to this theory, there are two perceptual networks: (1) the ventral stream responsible for object and face recognition, which is retained in patients with WS; and (2) the dorsal stream associated with object location and spatial management (Atkinson, 2017).

Several studies have provided evidence suggesting that the dorsal stream is related to occipito-parietal networks, specifically the intraparietal sulcus, precuneus, and inferior and superior parietal gyrus (Kravitz, Saleem, Baker, & Mishkin, 2011; Meyer-Linderberg et al., 2004). Global ability and closure have also been reported to be related to intraparietal sulcus alterations (Atkinson, 2017; Meyer-Linderberg et al., 2004; Mobbs et al., 2007). Our finding of impaired visual closure in WS patients without *GTF2IRD2* are consistent with prior findings showing that, like patients with right parietal lobe lesions, WS patients can identify and copy the elements of visual stimuli, but fail to integrate them into a whole (global process) (Atkinson et al., 2001; Atkinson & Braddick, 2012; Atkinson & Nardini, 2008). Hence, dorsal stream vulnerability theory (Atkinson & Nardini, 2008) could explain the greater visuospatial alterations in people with WS associated with loss of *GTF2IRD2*, relative to typical WS patients.

In this context, it is noteworthy that both the parietal cortex and cerebellum, areas with unusually high *GTF2IRD2* expression (Allen Brain Atlas, 2010; Porter et al., 2012; Uhlén et al., 2015), are involved in visuospatial ability and are altered markedly in WS (Martens, Wilson, & Reutens, 2008). The parietal and prefrontal cortices in the right hemisphere have been associated with the recognition of emotions in faces, judgment about them, theory of mind, and empathy (Adolphs, 2009; Shamay-Tsoory, Tomer, Goldsher, Berger,

& Aharon-Peretz, 2004). Structural and functional alterations have been found in the amygdala and right orbital-medial prefrontal cortex of WS patients performing tasks that evaluate these domains of social cognition (Bellugi et al., 2007; Mimura et al., 2010). WS patients perform similarly to patients with right parietal damage on visuospatial tasks (Atkinson et al., 2001; Atkinson & Braddick, 2012; Atkinson & Nardini, 2008); this greater deficit may be associated with lower visuospatial and social performance in those patients with a *GTF2IRD2* deletion.

Regarding behavioral profile, our finding that patients with a *GTF2IRD2* deletion presented several behavioral problems associated with social cognition indicate considerable challenges for these patients in appropriate expression of empathy, recognition of the emotional state of another, and theory of mind. This finding is consistent with previous results of Porter et al. (2012), who identified more difficulties in theory of mind tasks in patients with lost, versus retained, *GTF2IRD2*. Likewise, individual case reports of WS patients with co-deletion of *GTF2I*, *GTF2IRD1*, and *GTF2IRD2* described predominant autistic features (Edelmann et al., 2007; Karmiloff-Smith et al., 2012). Conversely, our adaptive skills scale results may indicate better social abilities performance and leisure activities in WS patients with conserved *GTF2IRD2*, and their relatively superior performance in these areas may be related to the preservation of their ability to perceive others' emotions and their more typical performance on tests of theory of mind and empathy.

Limitations

It is important to emphasize that in this study we did not apply specific or experimental tests that assessed the patients' cognitive-emotional abilities directly. However, the results of the inventories completed by the parents had moderate to large effect sizes. The limitations of this study include: (1) a lack of neuroimaging studies, which could identify structural inter-hemispheric differences (especially in the parietal and frontal cortices and in the cerebellum) associated with visuospatial processes and social cognition; (2) a lack of psychological and experimental tests that assess the components of social cognition directly; and (3) a small sample, which precludes a strong statistical analysis and limits the generalizability of the results. Notwithstanding, this work provides an approach to examining the relationship among genes, brain, cognition, and behavior in WS patients.

CONCLUSION

The present findings provide new evidence of the importance of *GTF2IRD2* in the cognitive, behavioral, and adaptive phenotypes of WS patients. The results support the notion that the locally elevated expression of *GTF2IRD2* in the right prefrontal and parietal cortices, when the gene is present,

reflect its role in the cognitive, behavioral, and adaptive functions that are impaired in WS patients with deletion of *GTF2IRD2*. Furthermore, given *GTF2IRD2*'s regulatory influence on *GTF2IRD1*, loss of *GTF2IRD2* may result in structural alterations in sites where *GTF2IRD1* is normally expressed, leading to more profound effects on visuospatial and social skills. It is necessary to continue studying the phenotypic outcomes associated with different deletion variants in WS patients to identify the neuropsychological influence of specific genes affected in WS.

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SUPPLEMENTARY MATERIALS

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