

Association Between The SNP Rs2154545 Polymorphism on The DyrkIA Gene and Intellectual Disability in Down Syndrome

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Abstract. Down syndrome, or trisomy 21, is a genetic abnormality caused when chromosome 21 fails to separate during meiotic division, resulting in offspring with 47 chromosomes. People with this syndrome are known to have mild to severe intellectual disability. One of the pivotal genes on chromosome 21 is Dual-specificity tyrosine-phosphorylation regulated kinase 1A (DyrkIA), and some of its polymorphisms (for example, SNP rs2154545) may cause phenotypic changes in a person with Down syndrome. The purpose of this study is to demonstrate whether SNP rs2154545 polymorphisms are associated with the DyrkIA gene and to analyze its relationship with intellectual disability in Down syndrome. This was a cross-sectional observational study that used blood samples from 39 subjects. All subjects were identified through Special School Kodya Padang, they were clinically proven to have Down syndrome, and the trisomy 21 diagnosis was confirmed by karyotyping analysis. Their DNA samples were analyzed using restriction fragment length polymorphism (RFLP), the samples were sequenced, and the contigs were qualitatively analyzed. Multiple alignment and BLAST-NCBI was used to confirm the restriction result. The rs2154545 polymorphism on the DyrkIA gene was observed in the form of homozygote mutant alleles (AA) in 5.1% of all samples and heterozygote mutant alleles (GA) in 51.3% of all samples. The remaining samples were wild-type (43.6%). Statistical analysis was performed, but there were no significant relationships ($p > 0.05$) between gene polymorphism and intellectual disability. The rs2154545 polymorphism on the DyrkIA gene was proven to exist in some people with Down syndrome, but there were no significant relationships between the polymorphism and every category of intellectual disability (that is, mild, borderline, moderate, and severe).

Keywords: Down Syndrome, DyrkIA Gene, Intellectual Disability, RS2154545 Polymorphism.

1 Introduction

Down syndrome (DS) is a major pediatric issue due to its impact on growth and development. Cognitive development is the most inhibited factor and causes hindrance in everyday life. People with DS need help and counseling from family, the community, and the country. DS is a genetic abnormality that result intellectual disability, hypotonia, facial dysmorphic, earlier onset of Alzheimer's disease, and other behavior disruptions. The incidence

tests in school-age children are considered to be a better reflection of the child long-term cognitive abilities [6].

Recent studies observed that DyrkIA plays a role in the interference of early cognitive development and brain degeneration, which causes the intelligence disruption as reflected by IQ scores. The present study has been performed to elucidate the DyrkIA gene polymorphism SNP rs2154545 and its relationship to intelligence in DS. The objectives of this study are to demonstrate whether the SNP rs2154545 polymorphism occurs in the DyrkIA gene and to analyze its relationship with intellectual disability in DS.

2 Materials and Methods

This study was conducted using blood samples from 39 students from the Special School Kodya Padang who meet the inclusion criteria. Up to 4 cc of blood was taken from each student's vein for cytogenetic analysis and DNA extraction. Polymerase chain reaction (PCR) and sequencing were performed in a biomedical laboratory in the medical faculty at Andalas University, Padang and Macrogen, South Korea. A professional psychologist performed IQ testing.

2.1 DNA Extraction (Genomic DNA Purification)

Genomic DNA was isolated from 300 µl blood sample using the Genomic DNA Mini Kit (Blood/Cultured Cell) GB100 (Geneaid). DNA isolation was performed by following the instructions in the kit manual, including sample preparation, cell lysis, DNA binding, washing and rinsing, and DNA elution.

2.2 Rs2154545 (G>A) Genotyping

Genotyping for the SNP Rs2154545 polymorphism on the DyrkIA gene (NCBI accession number: NG_009366.1) was performed using the restriction fragment length polymorphism (RFLP)-PCR method. PCR reagents consisted of: (1) 0.2 µM deoxynucleoside triphosphates (dNTPs); (2) 100 ng genomic DNA; (3) 1.25 U HotStarTaq DNA Polymerase with its buffer; (4) 0.5 µM forward primer rs2154545-F (5'-GGATTCTGTAGTGGATATGTAGTGG-3'); and (5) 0.5 µM reverse primer rs2154545-R

(5'-ACCCTGTTCCACGCTCAAAA-3'). All reagents were combined to create a total volume of 25 µl.

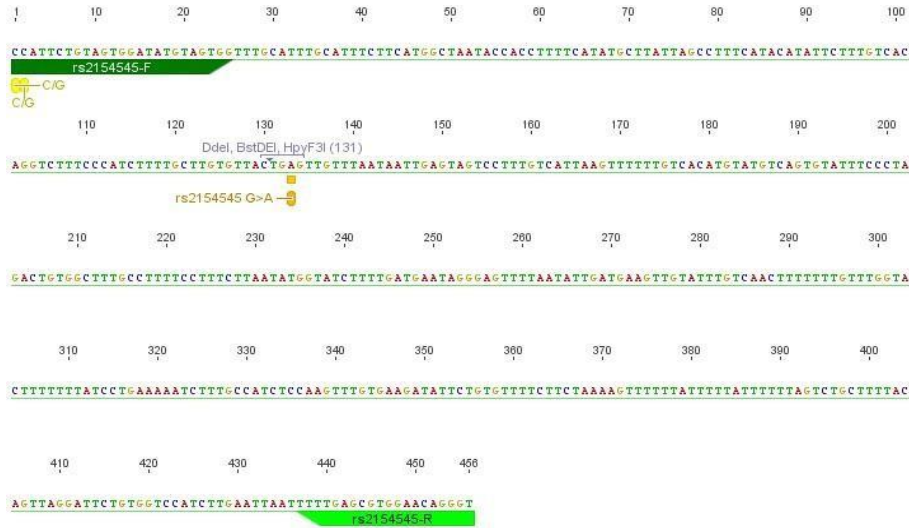


Fig. 1. RFLP-PCR map of DYRK1A (NCBI accession number: NG_009366.1) for rs2154545 (G>A) mutant allele with DdeI restriction site.

PCR was performed under the following conditions: (1) initial denaturation at 95°C for 5 minutes; (2) 35 cycles of denaturation at 94°C for 1 minute, annealing at 53°C for 30 seconds, and elongation at 72°C for 30 seconds; and (3) final elongation at 72°C for 10 minutes. The PCR product was examined using 1.5% agarose gel electrophoresis with DNA GelRed stain in GelDoc. The Rs2154545 PCR product was 456 bp in length.

The PCR product was cut by 1 U of DdeI restriction enzyme (restriction site= C[^]TNAG) at 65°C for 2 hours. After that, the restriction result was examined using 1.5% agarose gel electrophoresis with GelRed DNA stain in GelDoc. Samples from individuals with the wild-type rs2154545 genotype (GG allele) resulted in a DNA band that was 456 bp in length. In contrast, samples from individuals with the mutant heterozygote rs2154545 genotype (GA allele) produced 456 bp, 326 bp, and 130 bp DNA bands, and samples from individuals with the mutant homozygote rs2154545 genotype (AA allele) produced 326 bp and 130 bp DNA bands.

Results

Every subject in this study had trisomy 21, confirmed by karyotyping analysis. Of the 39 subjects, 64.1% (n=25) were male, 71.8% were greater than or equal to 10 years old, 74.4% were born from a mother who was greater than or equal to 35 years old, and 63.1% had mild–borderline intellectual disabilities (Table 1).

Table 1. Study Subjects Characteristics.

Characteristic	f=39	%
1. Gender		
* Male Trisomy	25	64.1
* Female Trisomy	14	35.9
2. Age		
≤ 10-year-old	11	28.2
> 10-year-old	28	71.8
3. Mother's age		
≤ 35-year-old	10	25.6
> 35-year-old	29	74.4
4. Nutrition status		
* Underweight	13	43.6
* Norm weight	18	30.8
* Overweight	8	25.6

3.1 Karyotyping Analysis

Blood samples were collected from all subjects and placed in a heparin-coated vacutainer and cultured during an incubation period of 3 to 4 days. The cells were then harvested and the harvested products were deposited onto a glass slide and left for 1 to 3 days. After that, the banding procedure was carried out. The banding result was analyzed and six metaphase cells were selected for karyotyping. In total, 47 chromosomes, XY, trisomy 21 (male) and 47 chromosomes, XX, trisomy 21 (female) were acquired (Figure 2).

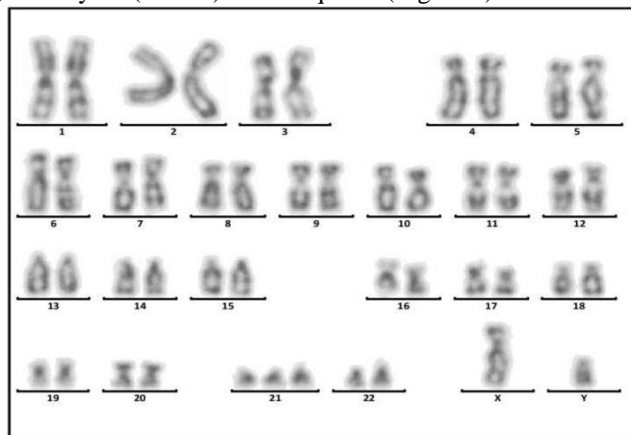


Fig. 2. DS 01 karyotyping result with 3 copies of 21st chromosome and XY sex chromosome which implies that DS01 patient is a male.

3.2 DNA Extraction (Genomic DNA Purification)

DNA extraction was carried out by following the instruction manual from Genomic DNA Mini Kit (Geneaid), and the sample was examined by gel electrophoresis. Every sample produced a DNA band (Figure 3), which implies that the genomic DNA was successfully isolated.



Fig. 3. Electrophoresis result of Isolated DNA from DS1 to DS20 patients. PCR Genotyping rs2154545 (G>A) Result.

After completion of the PCR and restriction enzyme procedures, DNA samples from four people (DS32, DS15, DS39, and DS31) were separated into three DNA bands — 456 bp, 326 bp, and 130 bp — which implies that the GA allele (mutant heterozygote) was present in those subject. The DNA sample from DS40 DNA was separated into two bands — 326 bp and 130 bp — which implies the AA allele (mutant homozygote). The DNA samples from DS26, DS27, and DS11 only produced one band (456 bp), which implies the GG wild-type allele (Figure 5).

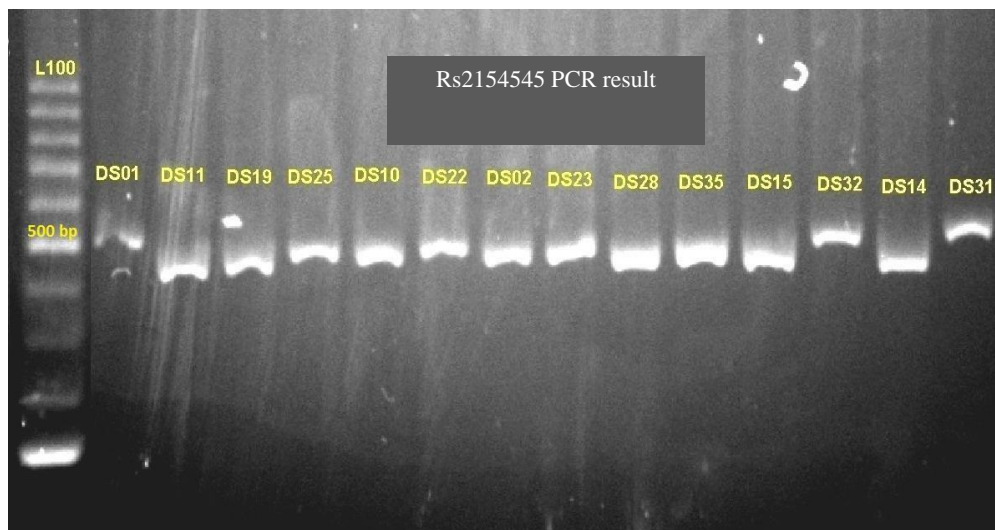


Fig. 4. Electropherogram of DS subject's rs2154545 PCR. Rs2154545 (G>A) PCR product was 456 bp band.

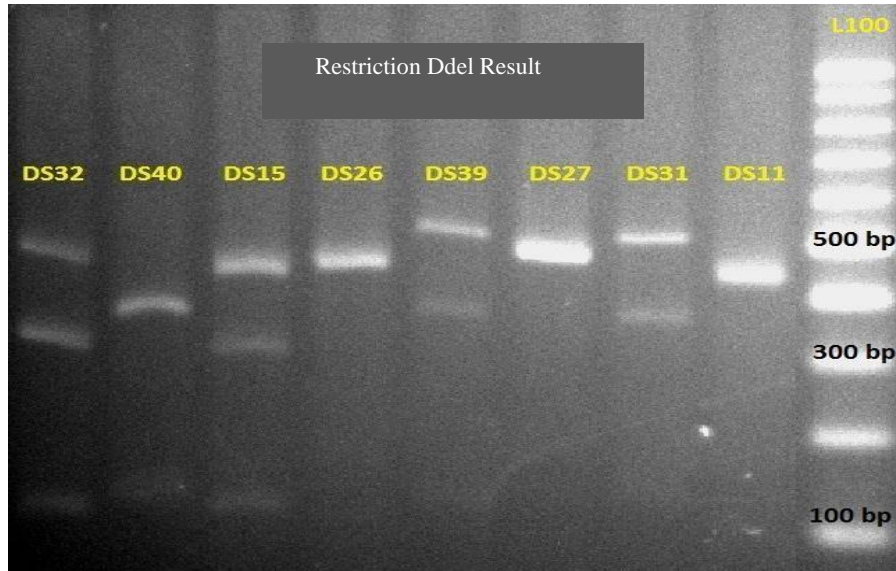


Fig. 5. Electropherogram of Ddel restriction enzyme cut of DS subject's rs2154545. Rs2154545 (G>A) PCR product was 456 bp band.

3.3 Sequencing

The PCR product from a few samples was purified and sent to Macrogen, South Korea, for sequencing. Below is the sequencing result of DS22.

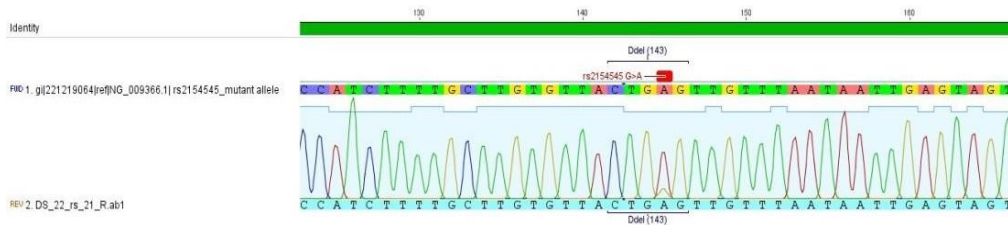


Fig. 6. Contig process of sequencing product of DS 22 DNA sample with forwarding primer for rs2154545 (G>A) SNP. Sequencing result indicated that DS22 is a mutant heterozygote (GA allele) which in agreement with the RFLP result.

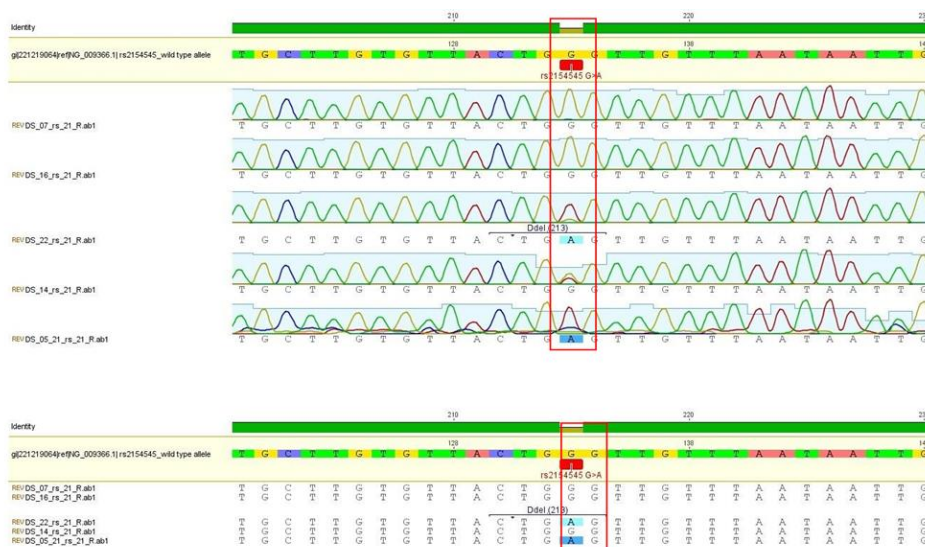


Fig. 7. Multiple alignments from 5 contigs of sequenced samples. Multiple alignments showed that DS 22 and DS 05 have SNP polymorphism rs2154545 in the Dyrk1A exon sequence as indicated by nucleotide substitution from G to An in DS 22 and DS 05.

3.4 Multiple alignment results

Of all 39 subjects sampled in this study, as many as 17 subjects (43.6%) had the wild-type allele (GG), two subjects (5.1%) were mutant homozygotes (AA allele), and 20 subjects (51.3%) were mutant heterozygotes (GA allele) (Table 2). Subjects with the homozygote mutant allele (AA) were equally distributed in the intellectual disability categories of mild–borderline and moderate-severe, but the results were not statistically significant ($p > 0.05$) (Table 3).

Table 2. Dyrk1A gene SNP rs2154545 Polymorphism

SNP rs2154545 Polymorphism	F	%
Wild Type (GG allele)	17	43.6
Mutant Homozygote (AA allele)	2	5.1
Mutant Heterozygote (GA allele)	20	51.3

Table 2 has heterozygote mutant 51.3 %

Table 3. Dyrk1A gene polymorphism rs2154545 relationship with Mental Retardation level

SNP rs2154545	Mental retardation				f	Total %	P
	Mild-Borderline		moderate-severe				
	f	%	F	%			
Wild Type	9	52.9	8	47.1%	17	100	0.355
Homozygote	1	50	1	50	2	100	
Heterozygot	15	74	5	25	20	100	
Total	25	64.1	14	35.9	39	100	

Table 3. has heterozygote mutant relationship with the mild borderline of mental retardation

4 Discussion

This study included 39 subjects who have been diagnosed with DS (trisomy 21). The syndrome is caused when a gamete cell with an extra copy of chromosome 21 undergoes fertilization with a normal gamete cell, resulting in a zygote with 47 chromosomes (46XX + 21 and 46XY + 21). This abnormality occurred in 90%–95% of all observed cases [8][9]. In this study, 25 subjects (65.1%) were male and 14 subjects were female (35.9%). This proportion is in accordance with results reported by Johnson (2006) in which more male babies were born with DS compared to female babies. [10]

The prevalence of intellectual disabilities in people with DS is around 2–3% of all cases. Most people (85%) who have been diagnosed with DS have intellectual disabilities that are considered to be mild, while 0.3%–0.5% have intellectual disabilities that are considered to be severe [11]. Mild intellectual disabilities are likely to occur 7–10 times more frequently compared to moderate and severe intellectual disabilities. Although the physical appearance of DS is easily identified, intellectual disability is the most consistent identifier in DS, and the average IQ of many people with this diagnosis equal to 50 [10]. This study observed 64.1% people with mild–borderline intellectual disability and 35.9% with moderate-severe intellectual disability.

In the present study, we observed 16 males diagnosed with DS who had mild–borderline intellectual disabilities and 9 males with moderate-severe intellectual disabilities. In the females with DS, 10 were considered to have mild–borderline intellectual disabilities and 4 males with moderate-severe intellectual disabilities. Statistical analysis showed that there were no significant relationships between the severity of intellectual disability and gender ($p > 0.05$).

As much as 6 under 10-year-old subjects suffered mild-borderline intellectual disability and 5 subjects in the same age group had a moderate-severe intellectual disability. 20 subjects, older than 10-year-old, had a mild-borderline intellectual disability, while 8 subjects in the same age group had a moderate-severe intellectual disability. Statistical analysis didn't reveal any significant relationship between age and intellectual disability level ($p > 0.05$).

Based on a study conducted in 2000, people with intellectual disabilities were identified in 12 of every 1,000 school-aged children, or one in every 83 children [11]. Intellectual disability mainly occurs in older people. These previously reported statistics are consistent with the findings of our, in which 11 subjects (28.2%) who were less than 10 years old had an intellectual disability compared to 28 subjects (71.8%) who were greater than or equal to 10 years old. As previously stated, our study identified 29 people (74.4%) who had mothers who were 35 years or older when giving birth to their child with DS. According to other authors, DS cases occur in one of every 700 births from mothers who are 35 years old or older, and the prevalence of DS births increases as the mother's age increases [12][9]. A study in South Africa also found a similar result.

DS can also occur as a result of genetic mutation. Some two hypotheses explain the etiology of DS as the result of mutation: (1) the instability hypothesis that describes the loss of chromosome stability; and (2) the gene dose hypothesis that describes the overexpression of a particular gene. Our study identified the SNP rs2154545 polymorphism on the Dyrk1A gene. The wild-type form (GG allele) was observed in 17 samples (43.6%), the mutant homozygote form (AA allele) in 2 samples (5.1%), and the heterozygote mutant form in 20 samples (51.3%).

In general, there were 22 rs2154545 SNP mutants (56.4%) of the DyrkIA gene. This result implies that DyrkIA polymorphisms have a role in the pathogenesis of DS. DyrkIA overexpression has the potency to damage the brain and cause intellectual disability.

In humans, the DyrkIA gene is located on the HSA 21 loci, the long arm of the 21q22.3 chromosome; this region is the critical site of DS and is usually called DScritical region (DSCR). DSCR, which is comprised of 5.4 Mb and 30 genes, has been characterized by phenotype-genotype correlation studies using samples from people diagnosed with partial trisomy 21 [14][5]. Some studies have shown that DyrkIA gene has an influence on the development of the central nervous system, such as neuron proliferation, neurogenesis, differentiation, cell death, and synapse plasticity. The overexpression of this gene in the brain of a fetus with DS supports the hypothesis that DyrkIA plays a role in neurodevelopment and the cognitive deficits of people with DS [4]. In our study sample, people with DS who were heterozygous for the SNP rs2154545 polymorphism were mainly found to have mild–borderline intellectual disabilities (74.0%), while half of the sample who were homozygous for the polymorphism had mild–borderline intellectual disabilities and half had moderate-severe intellectual disabilities. However, no findings reached statistical significance.

A study of seniors with Alzheimer's disease in Madrid, Spain, found variations of the DyrkIA gene (SNP polymorphisms rs117001483, rs2835740, rs1137600, rs2154545, rs8132976, and rs2835762) that did not reach statistical significance [15]. A study in a Chinese population found a relationship between mild cognitive amnesic disturbance and SNP polymorphisms in three systems that relate to Alzheimer's pathogenesis. Those three systems were the amyloid cascade pathway, Tau metabolism, and cholesterol [16]

In conclusion, the results of our study show that there are an observable SNP rs2154545 polymorphism on the DyrkIA gene has heterozygote mutant (Allele GA) that causes the most common clinical manifestation of this overexpression is a mild–borderline intellectual disability in people with DS.

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