

ASSOCIATION OF SINGLE NUCLEOTIDE POLYMORPHISM IN CHROMOSOME 11 WITH AUTISM SPECTRUM DISORDER

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Communicated by Isaak Rashal

Several genetic loci in chromosomes 11 and 15 have recently been associated with non-syndromic autism spectrum disorder (ASD) in populations from North America and Europe. The aim of the present study was to investigate whether such an association exists in a Latvian population. Ninety-five patients with ASD in the age range 3–20 years (mean age 8 years, SD 3.18) participated in the study. The control group consisted of 161 healthy, non-related individuals without ASD randomly selected from the Latvian Genome Database. Four single nucleotide polymorphisms (SNPs) — rs11212733, SNP rs1394119, rs2421826, rs1454985 — were genotyped by the TaqMan method. Allele frequency differences between ASD patients and control subjects were compared for each SNP using a standard chi-square test with Bonferroni correction. The level of statistical significance was set at 0.05 for nominal association. Only the genetic marker rs11212733, localised on the long arm of chromosome 11 in locus 22.3, was found to be strongly associated with the ASD patient group (χ^2 6.982, P_{adjusted} 0.033, odds ratio 1.625). Our data demonstrating a significant relationship between the SNP rs11212733 and the development of ASD in a Latvian population suggest that it is not a population-specific relationship. Thus, future studies focusing on the DDX10 gene and related genetic loci are needed.

Key words: autism spectrum disorder (ASD), single nucleotide polymorphisms (SNPs), rs11212733, DDX10.

INTRODUCTION

Autism spectrum disorders (ASD) are serious early childhood neurodevelopmental disorders with unknown etiology and a rapid annual increase in prevalence. ASD are a broad phenotype including less severe disorders (Anonymous, 1994; Bailly *et al.*, 1996; Johnson *et al.*, 2007). ASD are clinically characterised by impaired social and communication skills, hyperactivity and attention deficit, stereotypic movements and interests, stereotypic rituals, emotional disturbances, and varying degrees of expressive and receptive language development disorders (Bailly *et al.*, 1996; Risch *et al.*, 1999).

ASD is not a disease but rather a syndrome that is characterised by a multifactorial type of inheritance. In some cases, it is one of the symptoms of monogenic or chromosomal pathology, and can also be a symptom of inherited metabolic disorders. In early childhood, the latter can manifest primarily with autistic behaviour, speech development delay or re-

gression, and mental retardation. This collection of symptoms should be classified as ASD (Gillberg *et al.*, 2006). The International Classification of Diseases (ICD-10) distinguishes between autism and ASD, which are subdivided into syndromic autism and non-syndromic or idiopathic autism (Anonymous, 1992; Lintas and Persico, 2009).

Linkage and association studies have identified potential ASD candidate genes in different chromosomes — the most significant ones being 2q, 7q, 15q and the X chromosome (Muhle *et al.*, 2004; Lintas and Persico, 2009). As several genetic loci have been implicated in ASD development, extensive studies have been conducted worldwide to detect potential candidate genes for ASD (Liu *et al.*, 2008; Cho *et al.*, 2011). Genome-wide linkage analyses of quantitative and categorical autism subphenotypes, carried out in North America and Europe, revealed association between ASD and locus 15.4-p15.3 in the short arm of chromosome 11 (Liu *et al.*, 2008). Furthermore, cytogenetic abnormalities at

locus 11-13 in the long arm of chromosome 15 have been reported to cause ASD. Population studies and case reports describe the duplication, deletions and inversions of this locus, as well as the phenotype of chromosome 15 in ASD patients (Muhle *et al.*, 2004; Johnson *et al.*, 2011).

The aim of this study was to investigate whether the association between reported ASD informative genetic markers in chromosomes 11q, 11p, and 15q and non-syndromic autism spectrum disorders exists in a Latvian population.

MATERIALS AND METHODS

Subjects. The Children's Psychiatry Clinic of the Children's University Hospital is one of four specialised centres in Latvia. Since 2006, 195 patients with possible ASD have attended for a consultation. The given patient group was primarily examined by a psychiatrist using ICD-10 diagnostic criteria, as well as standardised scales corresponding to each patient's age: The Checklist for Autism in Toddlers (the CHAT) (assessment in the first 18 months); Autism Spectrum Quotient (autism spectrum questions for children aged 4–11 years); Cambridge University Behaviour and Personality questions for children up to the age of four years; Cambridge University social and communicative development questions; and Childhood Asperger Syndrome Test (CAST) (Anonymous, 1992; Baron-Cohen *et al.*, 1992; Scott *et al.*, 2002; Baron-Cohen *et al.*, 2006; Williams *et al.*, 2006). In order to specify the diagnosis of ASD in cases with positive test results, they were referred to a psychologist for the Autism Diagnostic Observation Schedule (ADOS) test, using the most appropriate model of the given test for each patient (Lord *et al.*, 2002). Thus, on the basis of the test results, of the 195 patients, diagnoses of ASD were confirmed in 169 and were selected.

A clinical geneticist performed the genetic assessment of all patients. Seventy-four patients were excluded from the group as monogenic, chromosomal and metabolic pathologies, diagnosed using standard karyotype, long arm of chromosome 15 locus 11.13 (15q11.13) deletion/duplication and long arm of chromosome 22 locus 11.2 (22q11.2) deletion analyses.

Ninety-five patients with ASD in the age range 3–20 years (mean age 8 years, SD 3.18) agreed to participate in the study. The male to female ratio was 3.5:1. The patients, legal guardians or parents signed informed consent forms in accordance with the instructions issued by the Medical Ethics Committee of Riga Stradiņš University.

The control group comprised 161 healthy, non-related individuals randomly selected from the Latvian Genome Database. A subject was excluded from the control group if there was information concerning possible mental illness. The male to female ratio was 1.7 : 1.

Analysed patients and control group individuals corresponded to the ethnic structure of the general Latvian population.

Methods. DNA was extracted from each patient's venous blood sample (collected at the Children's Psychiatry and Medical Genetics Clinics of the Children's University Hospital) using a standard phenol/chloroform extraction protocol (Sambrook *et al.*, 1989).

Single Nucleotide Polymorphism (SNP) Selection. On the basis of the data of recent reports on the most significant SNPs involved in the development of ASD, four SNPs were selected for the genotyping assay: rs11212733 at the long arm of chromosome 11 in locus 22.3, SNP rs1394119 at the short arm of chromosome 11 in locus 15.4–15.3, rs2421826 at the short arm of chromosome 11 in locus 13 and rs1454985 at the long arm of chromosome 15 in locus 13.3–14 (Liu *et al.*, 2008; Cho *et al.*, 2011).

TaqMan Genotyping. SNPs were genotyped using TaqMan® probe-based chemistries (Applied Biosystems, Carlsbad, CA, USA) on an automatic sequence-detection instrument (Real-Time PCR System, Applied Biosystems). All reactions were carried out using standard conditions as recommended by the manufacturer.

Statistical analysis. Allele frequency differences between ASD patients and control subjects were compared for each SNP using a standard chi-square test with Bonferroni correction. Allelic odds ratios and 95% confidence intervals were estimated using a standard chi-square test, assuming a multiplicative model. The level of statistical significance was set at 0.05 for nominal association. Statistical analyses were conducted using PLINK 1.06 software (Purcell *et al.*, 2007).

RESULTS

Four SNPs in the analysed genetic loci related to ASD were genotyped in 95 patients with ASD and 161 healthy, non-related controls. The average genotype call rate for these SNPs was 99.8%.

The genotype distributions in the study groups were in Hardy-Weinberg equilibrium, except for rs1394119, which was excluded from further studies. The genomic control inflation factor (kGC) was 1.008 for the entire data set. The results are shown in Table I.

Our results showed that the genetic marker rs11212733, localised on the long arm of chromosome 11 in locus 22.3, was found to be associated with ASD ($\chi^2 = 6.982$, $P_{\text{adjusted}} = 0.033$, odds ratio = 1.625). The genetic markers rs2421826, localised on the short arm of chromosome 11 in locus 3 and rs1454985, localised on the long arm of chromosome 15 in locus 13.3–14 did not have an association with ASD ($\chi^2 = 0.554$, $P_{\text{adjusted}} = 1.0$, odds ratio = 1.154; $\chi^2 = 1.118$, $P_{\text{adjusted}} = 1.0$, odds ratio = 1.217, accordingly).

DISCUSSION

Although we did not detect an association of rs2421826 or rs1454985 with ASD (perhaps due to the small number and

Table 1

RESULTS OF CASE-CONTROL ASSOCIATION STUDY

Chr	SNP	MAF cases	MAF controls	χ^2	P value	P_{adjusted}	Odds ratio	95% CI
11q22.3	rs11212733	0.552	0.432	6.982	0.008	0.033	1.625	1.13–2.31
11p13	rs2421826	0.352	0.432	0.554	0.456	1.0	1.154	0.79–1.68
15q13.3–q14	rs1454985	0.447	0.399	1.118	0.291	1.0	1.217	0.84–1.75

Chr, chromosomal localisation; SNP, single nucleotide polymorphism; MAF, minor allele frequency; P_{adjusted} , Bonferroni correction for three markers applied; CI, confidence interval

more homogeneous nature of our non-syndromic ASD cases), the SNP rs11212733, in chromosome 11q22.3 and located to the 5' region of the exophilin 5 (*EXPH5*) gene and 3' of the DEAD (Asp-Glu-Ala-Asp) box polypeptide 10 (*DDX10*) gene, was found to be strongly associated with our ASD patient group ($\chi^2 = 6.982$, $P_{\text{adjusted}} = 0.033$, odds ratio = 1.625, 95% CI = 1.13–2.31). This genetic marker has recently been reported to be significant in a Korean population (Cho *et al.*, 2011). As our Latvian ASD patients belong to a North-Eastern European population, which is distinct from a Korean population (Asian), our data suggest that the genetic marker at locus 11q22.3 is linked to a genetic locus that plays a prominent role in the development of ASD and is not population-specific.

EXPH5 is a protein-coding gene localised at the long arm of chromosome 11 in locus 22.3. A known function of this gene is related to Rab27, which governs the regulated exocytosis in non-neuronal cells, such as lytic granule secretion in cytotoxic T cells, production of insulin in pancreatic β -cells, as well as the release of histamine-containing granules in mast cells. Rab27 is also responsible for melanosome transport along the actin cytoskeleton in melanocytes (Kondo *et al.*, 2006). As *EXPH5* does not appear to be involved in neurological processes, it is probably not of notable significance in ASD.

The *DDX10* gene at the long arm of chromosome 11 in locus 22–23 encodes an RNA helicase, involved in RNA duplex unwinding and ribosome assembly. Although it is not known which RNA molecules are unfolded by this helicase, impaired function of *DDX10* would be expected to cause regulatory problems at the RNA level (Savitsky *et al.*, 1996). Thus, *DDX10* is a more promising candidate gene for further research concerning the development of ASD.

Our data demonstrating a significant relationship between the SNP rs11212733 and the development of ASD in a Latvian population suggest that it is not a population-specific relationship. Thus, future studies focusing on the *DDX10* gene and related genetic loci are needed.

ACKNOWLEDGMENTS

This study was supported by the European Social Foundation (ESF), Project No. 2009/0147/IDP/1.1.2.1.2/09/IPIA/VIAA/009. We are grateful to all the parents and children for their participation in this study.

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Received 21 November 2012

AUTISKĀ SPEKTRA TRAUCĒJUMU SAISTĪBA AR 11. HROMOSOMAS GENĒTISKO POLIMORFISMU

Autiskā spektra traucējumi (AST) ir agrīns, komplekss psihiskās attīstības bojājums, kurš sākas un tiek diagnosticēts bērniem līdz trīs gadu vecumam. AST ir multifaktoriāla ģenētiska slimība, kuras izcelsmē iesaistīti daudzi iespējamie kandidātgēni. Veikto gadījuma kontroles asociācijas pētījumu mērķis bija noskaidrot vai pastāv 11. un 15. hromosomu ģenētiskā polimorfisma saistība ar AST Latvijas populācijā. Balstoties uz literatūras datiem, tika izvēlēti četri potenciāli nozīmīgi SNP (rs11212733, rs1394119, rs2421826, rs1454985), kuri varētu būt iesaistīti AST attīstībā. Pētāmo grupu veidoja VSIA BKUS BS Gailezers Bērnu psihiatrijas un Medicīniskās ģenētikas klinikas 95 pacienti ar autismu un AST, kuru vecāki vai aizbildni rakstiski piekrita bērna ģenētiskā materiāla tālākai izpētei. Pētījuma kontroles grupā iekļāva nejaušināti atlasītas 161 potenciāli veselas, savstarpēji neradniecīgas personas no Valsts iedzīvotāju genoma datubāzes, kuriem anamnēzē nav datu par AST vai citiem psihiskiem traucējumiem ģimenē. Izvēlēto alēlu genotipēšana veikta Latvijas Biomedicīnas Pētījumu un studiju centrā, izmantojot *TaqMan* reaktīvus. Statistiskā datu analīze veikta, izmantojot PLINK 1.06 programmatūru, salīdzinot AST un kontroles grupu. Datu statistiskai apstrādei izmantots standarta $H\bar{I}$ kvadrāta (χ^2) tests ar Bonferoni korekciju. Pētījumā atklāta statistiski ticama SNP rs11212733 (lokalizēts 11. hromosomas garā plecā 22.3 lokusā) saistība ar AST attīstību Latvijas populācijā ($\chi^2 = 6,982$; P koriģēts $0,033$; izredžu attiecība $1,625$). Literatūrā aprakstītais polimorfisms ir saistīts ar autiskā spektra traucējumiem Dienvidkorejas pacientiem. Rs11212733 atrodas 11. hromosomas nekodējošā rajonā starp gēniem EXPH5 un DDX10, kuru nozīme autisma attīstībā vēl nav pierādīta. Nepieciešami turpmāki pētījumi.