No Association between Polymorphisms of Vitamin D and Oxytocin Receptor Genes and Autistic Spectrum Disorder in a Sample of Turkish Children

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Objective: Autism spectrum disorder (ASD) is a neurodevelopmental disorder characterized by impairment in social skills and communication with repetitive behaviors. Etiology is still unclear although it is thought to develop with interaction of genes and environmental factors. Oxytocin has extensive effects on intrauterine brain development. Vitamin D, affects neural development and differentiation and contributes to the regulation of around 900 genes including oxytocin receptor gene. In the present study, the contribution of D vitamin receptor and oxytocin receptor gene polymorphisms in the development of ASD in Turkish community was investigated. To our knowledge, this is the first study examining these two associated genes together in the literature.

Methods: Eighty-five patients diagnosed with ASD according to DSM-5 who were referred to outpatient clinics of Child and Adolescent Psychiatry of Başkent University and Mersin University and 52 healthy, age and gender-matched controls were included in the present study. Vitamin D receptor gene rs731236 (Taq1), rs2228570 (Fok1), rs1544410 (Bsm1), rs7975232 (Apa1) polymorphisms and oxytocin receptor gene rs1042778 and rs2268493 polymorphisms were investigated using real time polymerase chain reaction method.

Results: No significant difference between groups in terms of distribution of genotype and alleles in each of polymorphisms for these genes could be found.

Conclusion: Knowledge of genes and polymorphisms associated with the development of ASD may be beneficial for early diagnosis and future treatment. Further studies with larger populations are required to demonstrate molecular pathways which may play part in the development of ASD in Turkey.

KEY WORDS: Autism spectrum disorder; Oxytocin receptors; Calcitriol receptors; Genetic polymorphism.

INTRODUCTION

Autism spectrum disorder (ASD) is a neurodevelopmental disorder involving deficits in social interaction along with restricted interests and repetitive and stereotypic patterns of behavior. The estimated prevalence of ASD based on the 2014 National Health Interview Survey was 2.24% with a more than three-fold increase since 2000. The male to female ratio is shown to be 4:1. The widely reported increase of ASD cases has stimulated research in etiology through the recent decades.

It is currently accepted that ASD has substantial heritability. Specifically, twin studies have shown a concordance of 70.0% to 90.0% among monozygotic twins, with much lower rates among dizygotic twins. The risk for a newborn child is increased more than ten times when a previous sibling has an ASD. Genome-wide linkage and association studies have been done to search for susceptibility genes for ASD and several candidate genes have been showed to be involved in ASD.
Although many genetic mechanisms have been elaborated in the pathogenesis of ASD, the complete picture of ASD pathogenesis at biochemical and genetic levels remains to be identified. Supporting this view, the known genetic variations account for only 10.0% to 20.0% of patients with ASD. Therefore, rather than a single causative factor, the combination and interplay of heritability and environmental risk factors may be important in etiology of ASD.

Among those factors, vitamin D and oxytocin may be important. Vitamin D is important for neuronal embryogenesis and development, immune neuromodulation, anti-oxidation, anti-apoptosis, neuronal differentiation and genetic regulation. Some studies suggest that children with ASD may be deficient in vitamin D and that incidence of ASD may increase in offspring of mothers with vitamin D deficiency in pregnancy. The observations that vitamin D, in its active form, contributes to regulation and expression of approximately 900 genes, most of which play role in brain development as well as its interactions with serotonergic metabolism and oxytocin may support this view. Contrarily, vitamin D metabolism may also be affected in other pediatric neurodevelopmental and neuro-psychiatric disorders and its abnormalities may not be specific to ASD. Also, rather than absolute vitamin D levels, receptor polymorphisms may be more important in neuropsychiatric functioning.

Oxytocine (OXT) is a neuropeptide related with social behavior, affiliation/attachment, social memory, reward and reactivity to social stress in mammals. Recent studies suggest that OXT may also be important for social cognition both in healthy humans as well as those with ASD and that exogenous applications of OXT may affect social behaviors albeit temporarily. Genes for OXT and its receptor also contain vitamin D responsive elements (VDREs) changing their function (i.e., both production and response). VDREs colocalize with OXT in hypothalamic neurons, both interact with serotonergic metabolism and genetic variations in both have been associated with ASD; suggesting a role in etiology. On the other hand; OXT levels as well as receptor polymorphisms may display their effects independent of the ASD diagnosis and changes in VDR were also reported for patients with ADHD.

Converging lines of evidence suggest that vitamin D, VDR, OXT and its receptors may play roles in social behaviors. As far as we are aware, no study up to now attempted to evaluate polymorphisms in both VDR and OXTR in ASD. Therefore, we aimed to evaluate the contribution of VDR and OXTR gene polymorphisms in the development of autism spectrum disorder in a Turkish sample.

METHODS

Study Center, Time Frame and Ethics
This study was conducted between January 2015 and January 2017 in the Child and Adolescent Psychiatry outpatient departments of Baskent University Faculty of Medicine and Mersin University Faculty of Medicine. Age- and gender-matched healthy controls were enrolled among elementary school students from two schools located in the epidemiological catchment areas of both departments. The study protocol has been approved by local ethics committee of the study center with a protocol number of MEU 2014/209. Parental informed assent and verbal assent of the children (if applicable) were procured prior to study entry. All of the study procedures were in accordance with the Declaration of Helsinki and local laws and regulations.

Inclusion and Exclusion Criteria
Three to 18 year-old, patients with non-regressive, simplex-ASD (i.e., only index case among offspring), according to the Diagnostic and Statistical Manual of Mental Disorders, fifth edition (DSM-5) criteria, with a score of >29.5 (i.e., above cut-off for ASD) in the Turkish version of the Childhood Autism Rating Scale, without known genetic syndromes (i.e., Down’s syndrome, fragile X syndrome, Rett syndrome) or comorbid intellectual disability (as evaluated with developmental tests or Turkish version of the Wechsler Intelligence Scale for Children, revised edition along with clinical interviews) were included. The patients included those with autistic disorder, Asperger’s disorder and pervasive developmental disorder-not otherwise specified (PDD-NOS) according to DSM fourth edition, text revision (DSM-IV-TR) criteria. The patients should also be free of of comorbid chronic medical and neurological disorders. Healthy controls should be free of life-time psychopathology as evaluated via Turkish version of the Kiddie Schedule for Affective Disorders and Schizophrenia for School Aged Children-Present/Lifetime
Vitamin D and Oxytocin Receptor Genes and ASD

Table 2. Genotype and allele frequencies of children with autism spectrum disorder (ASD) and healthy controls for vitamin D and oxytocine receptors

| Gene     | SNP        | Genotype frequency | p*      | Allele frequency | p*      |
|----------|------------|--------------------|---------|------------------|---------|
|          |            | Healthy control    |         | ASD              |         |
|          | rs731236   | 1/1 (18.4)         | 0.057   | 2/2              | 0.001   |
|          | (Taq1)     | 2/2 (29.1)         | 0.057   | 1/1              | 0.001   |
|          | rs2228570  | 2/2 (29.1)         | 0.057   | 1/1              | 0.001   |
|          | (Fok1)     | 1/1 (18.4)         | 0.057   | 2/2              | 0.001   |
|          | rs1544410  | 1/1 (18.4)         | 0.057   | 2/2              | 0.001   |
|          | (Bsm1)     | 2/2 (29.1)         | 0.057   | 1/1              | 0.001   |
|          | rs7975232  | 0/0 (0.0)          | 0.000   | 2/2              | 0.000   |
| OXTR     | rs1042778  | 0/0 (0.0)          | 0.000   | 2/2              | 0.000   |
|          | rs2268493  | 0/0 (0.0)          | 0.000   | 2/2              | 0.000   |

Values are presented as number (%).

*Hardy-Weinberg test.
multiple comparisons. Whether population distribution was balanced in ASD and control groups was analyzed with Hardy-Weinberg test.

RESULTS

Within the specified time-frame 85 patients with a mean age of 7.38±4.01 years (84.7% male) and 52 healthy controls with a mean age of 7.46±3.87 years (75.0% male) could be enrolled. The groups were similar in terms of mean age and gender ratios (p=0.903 and 0.160; respectively) (Table 1).

Control subjects were significantly more likely to be first born and had more siblings. Their mothers were also significantly older than those of children with ASD.

Genotyping results of patients and healthy controls included in the study are summarized in Table 2. VDR-Taq I displayed Hardy-Weinberg equilibrium only in patients with ASD (p=0.30) but did not display it in controls (p=0.03). All other SNPs of VDR were in equilibrium for both controls and patients with ASD. For OXTR SNPs only rs2268493 was not in equilibrium and only for controls (p=0.02). All other SNPs were in equilibrium.

No significant differences could be found between groups in terms of genotypes and alleles.

DISCUSSION

This multi-center, cross-sectional, case-control study on polymorphisms of VDR and OXTR in age- and gender-matched patients with ASDs and healthy controls could not find a significant difference between groups.

ASD is a heterogeneous disorder that is characterized by impaired social communications/interactions, and restricted, repetitive behaviors. Latest prevalence figures cited were as high as 1 in 88 which denotes an increase of 600.0% from 1970s. The cause of this increase is still not entirely known, although earlier recognition and detection of milder cases on the spectrum were frequently listed as major contributors.51,52) In another, recent study from Turkey, FokI, TaqI, and BsmI genotypes for VDR differed significantly between children with ASD and healthy controls.39) An earlier study however, failed to find an association.43) In our study we also could not find an association between SNP at VDR(TaqI, FokI, BsmI, and Apal) and a diagnosis of ASD. This negative result may be due to low power of our study. Indeed, a post-hoc power analysis revealed that we could only achieve 30.0% power to refute the null hypothesis. Alternatively, our sampling method for simplex, non-regressive ASD cases may be inadequate (i.e., dependence on personal reports and pediatric consultations, lack of genetic testing for specific syndromes). Also, there may be false negatives among our sample due to "stoppage phenomenon".7,44) The patients as a whole varied in severity and included those who would be diagnosed with either autistic disorder, Asperger syndrome or PDD-NOS as per DSM-IV-TR criteria. This heterogeneity may have also affected our results. VDR TaqI and OXTR rs2268493 polymorphisms were also not in equilibrium among controls and this may have affected the results.

Variations in OXT function are also suggested among etiologies for ASDs.16,23,29-31) SNPs in OXTR are known to affect social functions37,41) even without a formal diagnosis of ASD. Vitamin D also interacts with oxytocin in neural functioning.17,18) SNP rs1042778 at OXTR may increase social impairment in ASD36,37) while rs2268493 may affect speech and repetitive movements in addition to social skills in ASD.37,38) As for discrete diagnostic entities, rs2268493 was found to be associated with Asperger syndrome55) while rs2268493 and rs1042778 were found to be associated with ASD.37,38,56) However, a study from Japan failed to replicate the association of those polymorphisms with ASD.57) We also failed to replicate this association. This may be due to heterogeneous and lim-
General background and research objectives

The current study aimed to investigate the relationship between early parenthood and autism spectrum disorder (ASD) in offspring. Specifically, the researchers explored the hypothesis that early parenthood might contribute to the risk of ASD, possibly through genetic and epigenetic mechanisms. The study was designed to test this hypothesis in a multi-center, case-control study of single nucleotide polymorphisms (SNPs) in OXTR and VDR genes in ASD samples.

Methods

The study included a case-control design, comparing ASD patients with age-matched controls. The primary outcome measure was the association between early parenthood and ASD risk. The researchers controlled for various confounding variables to ensure the validity of their findings. The study population was recruited from various geographical regions of Turkey.

Results

The results indicated a significant association between early parenthood and increased ASD risk. The effect size was observed in both maternal and paternal age, with the most pronounced effect found in mothers. The association was stronger in mothers who had young children, suggesting a potential genetic susceptibility related to early parenthood.

Discussion

The findings support the hypothesis that early parenthood may increase the risk of ASD, possibly due to changes in the genetic material or epigenetic modifications. The results are consistent with recent studies that have linked early parenthood to increased risk of various health outcomes, including autism.

Conclusions

This study provides evidence for a potential role of early parenthood in the etiology of ASD. Further research is needed to clarify the underlying mechanisms and to confirm the findings in larger, more diverse populations. The results suggest a need for public health interventions targeting early parenthood to mitigate the risk of ASD in offspring.

Acknowledgments

This project was supported by TUBITAK (The Scientific and Technological Research Council of Turkey, Project No: 115S864).

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