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PARTPEN STUR		DY OF ABNORMAL PALMER CREASES IN LDREN WITH AUTISM SPECTRUM ORDER: A CASE-CONTROL STUDY IN RTH INDIA	KEY WORDS:			
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ABSTRACT	INTRODUCTION: A social and communic formation of minor phy AIM: To compare the developing children. METHODOLOGY: It of Pt. B.D. Sharma, PG group) and fifty typica used to click photogr spectrum disorder we RESULTS: The prevale prevalence of Sydney crease in case group w CONCLUSION: Child as autism spectrum di of parental stress and b	atism Spectrum Disorder is a neurodevelopmental disorder cation patterns. Aberrant gene environment interactions de viscal anomalies such as abnormal palmar creases commonly se prevalence of abnormal palmar creases in children with autis was a case controlled cross sectional study conducted in depar IMS Rohtak. Fifty children of age 4-16 years with diagnosis ally developing children (control group) were recruited. A di aphs of the palms of children. Palmar crease patterns of fifty re compared with the control group. ence of abnormal palmar creases in case group was higher (47 crease in case group was double (22%) as compared to c crease in case group was 21%, while in control group it was ras 4%, while it was not seen in control group. Iren with abnormal palmar creases help in early screening of m sorder helping in early management of these children leading purden.	characterized mainly by deficits in luring fetal development leads to seen in autism spectrum disorder. sm spectrum disorder and typically rtments of Psychiatry and Pediatrics of autism spectrum disorder (case igital camera of 13 megapixels was y children with diagnosis of autism '%) than in control group (14%). The one in control group i.e. 11%. The only 3%. The prevalence of Suwon neurodevelopmental disorders such g to better outcomes and alleviation			

INTRODUCTION

Autism Spectrum Disorder (ASD) is a heterogeneous neurodevelopmental disorder characterized mainly by deficits in social and communication patterns.[1] DSM-5 groups the symptoms of ASD into two main categories: a) deficiencies in social interaction and communication, and b) patterns of repetitive and stereotypic movements.[2] This disorder is the second most common developmental disability, after intellectual disability. [3,4] The etiology of autism spectrum disorder is a complex interaction of genetics with environmental insults, which may lead to an aberrant neurodevelopment.[5] This aberrant neurodevelopment may also be the cause of various minor physical anomalies commonly seen in children with ASD. As brain and the skin develop from the same neuroectodermal layer during the early fetal development, these minor physical anomalies can be the reflection of aberrant development of brain during embryogenesis.[6] There are various minor physical anomalies such as epicanthus, hypertelorism, adherent ear lobes, abnormal palmar creases and high steeped palate commonly seen in patients with autistic spectrum disorder.[7] Abnormal palmar creases are perhaps one of the first minor physical anomalies apparent in these patients. Palmar creases are formed well within the first trimester of pregnancy and there is no alteration thereafter. [8] Any insult at this stage of organogenesis would lead to altered brain development and abnormal palmar creases. So, presence of abnormal palmar crease may indicate the altered development of brain during early embryogenesis. The aim of the present study was to compare the prevalence of abnormal palmar creases in children with autism spectrum disorder and typically developing children.

MATERIALS AND METHODS

The index study was a case controlled cross sectional study conducted in the out-patient departments of Psychiatry and Pediatrics, of Pt.B.D.Sharma, PGIMS Rohtak. Ethical clearance was sought from Institutional Ethics Committee. Written informed consent from the parent and assent was taken from the children participating in the study.

Fifty children of 4-16 years of age of either gender, with diagnosis of autism spectrum disorder as per DSM-5 criteria, and their parents attending the Child Guidance Clinic at department of Psychiatry, Pt. B.D. Sharma PGIMS, Rohtak constituted the study (case) group. Children with severe and profound mental retardation were excluded from this group. Fifty typically developing children, age and gender matched with the case group attending the Out-patient department of Pediatrics, Pt. B.D. Sharma PGIMS, Rohtak constituted the control group. The children were with major congenital malformations were excluded from this group.

After initial screening, final diagnosis of autism spectrum disorder was confirmed by the consultant psychiatrist. A specially designed proforma was used to gather sociodemographic and clinical details about the children and their parents. A digital camera of 13 megapixels (resolution of 3120 X 4160) was used to click photographs of the palms of children in both groups, with white sheet of paper as a clear background. Camera lens was kept perpendicular to the line joining distal flexion crease of wrist and proximal flexion crease of the middle finger while clicking the photographs.[9] Photographs were evaluated in both the groups for palmar crease patterns. Children with diagnosis of autism were further assessed with Childhood Autism Rating Scale (CARS) and Indian Scale for Assessment of Autism (ISAA). [10,11]

The data thus collected was analyzed using descriptive statistics for discrete and continuous variables. Chi-square test was used for comparing the categorical variables and ttest was used for comparing the continuous variables.

RESULTS

There were total of one hundred children, fifty children in case and control group each. Table 1 shows various socio-

demographic variables in both the groups. Majority of children in both groups were males and of age range of 4 to 7 years. Majority of the case group children had completed their primary education only as compared to the control group wherein children had moved to higher classes. In case group, 56% children belonged to the urban background while majority of the children in control group belonged to the rural background.

TABLE 1 Various socio-demographic variables of case and control groups

Variables	Cases	Controls	t/x2	p -value
	(n=50)	(n=50)		
Age				1 000
4-7 years	20	20	0 000	Not
8-11 years	13	13	0.000	significant
12-16 years	17	17		significant
Gender				1.000
Male	43	43	0.000	Not
Female	07	07		significant
Educational status				
Primary (up to 5th	44	32	07 00	0.007
standard)			21.39	0.005
Secondary (6th-	06	18	3	Significant
10th)				
Birth order status				
1	28	21		0.539
2	15	18	2.965	Not
3	05	08		significant
>3	02	03		5
Number of				
children in family				0.104
1	10	4		0.124
2	26	23	1.155	Not
3	09	18		significant
>3	05	05		
Adoption status				
Yes	0	0	-	-
No	0	0		
Number of family				
members				0.500
3-5	26	29	8.241	Not
6-8	16	12		Significant
9-11	08	09		
Family structure				0.276
Nuclear	32	29	2 576	Not
Extended	06	12	2.510	significant
Joint	12	09		significant
Neighborhood				0.098
status			4 652	Not
Rural	22	32	7.002	significant
Urban	28	18		Significant

TABLE 2 Types of palmar crease both hands in case and control groups

Palmar crease	Cases (N=100 hands)		Controls (N=100 ha	P Value	
pattern (both hands)	Frequency	%	Frequenc y	%	
Normal	53	53	86	86	< 0.001
Simian	22	22	11	11	Significant
Sydney	21	21	3	3	
Suwon	4	4	0	0	
	100	100	100	100	

TABLE 3 Types of palmar crease right and left hand in case and control groups

Palmar	Right hand		Left hand	
crease	Cases	Controls	Cases	Controls
	(n=50)	(n=50)	(n=50)	(n=50)

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	Frequency (%)	Frequency (%)	Frequency (%)	Frequen cy (%)
Normal	29 (58)	43 (86)	24 (48)	43 (86)
Simian	12 (24)	5 (10)	10 (20)	6 (12)
Sydney	9 (18)	2 (4)	12 (24)	1 (2)
Suwon	0 (0)	0 (0)	4 (8)	0 (0)
P value	0.007 Significant		<0.001 Significant	

Table 4. Correlation of palmar crease patterns both hands of case group with gender

Palmar Crease	Males(86	Females(14	PValue
Pattern	hands)	hands)	
Normal	44 (51.1%)	09 (64.2%)	0.119
Simian	17 (19.7%)	05 (35.7%)	Not
Sydney	21 (24.4%)	00 (0%)	significant
Suwon	04 (4.6%)	00 (0)	

Table 5. Severity of autism in case group by Childhood Autism Rating Scale-2nd edition (CARS-2)

CARS-2 scores	Cases (n=50)		
	Frequency	%	
30-36.5 (mild to moderateautism)	36	72	
37-60 (severe autism)	14	28	
	50	100	

Table 6 (a). Correlation of abnormal palmar creases (right hand) with scores on Childhood Autism Rating Scale-2nd edition(CARS-2)

Palmar crease	Cases	CARS-	2 scores	р
right hand	(n=50)	Mean	Std. Deviation	value
Normal	29	34.62	3.34	0.570
Simian	12	33.92	2.59	Not
Sydney	9	35.56	4.81	significant
	50	34.62	3.46	

Table 6 (b). Correlation of abnormal palmar creases (left hand) with scores on Childhood Autism Rating Scale- 2nd edition (CARS-2)

Palmar Cases		CARS-2	p value	
crease left hand	(n=50)	Mean	Std. Deviation	
Normal	24	34.13	3.61	0.714
Simian	10	34.50	2.49	Not
Sydney	12	35.50	3.76	significant
Suwon	4	35.25	4.35	
	50	34.62	3.46	

Table 7. Severity of autism in case group by Indian Scale for Assessment of Autism (ISAA)

ISAA Score	Cases (n=50)	
	Frequency	%
70-106 (mild)	15	30
107-153 (moderate)	27	54
>153 (severe)	08	16
	50	100

Table 8 (a) Correlation of abnormal palmar creases (right hand) with scores on Indian Scale for Assessment of Autism (ISAA)

Palmar creases right	Cases (n=50)	ISAA scores		p value
hand		Mean Std. Deviation		
Normal	29	116.14	27.99	0.140
Simian	12	104.75	24.07	Not
Sydney	9	129.44	31.00	significant
	50	115.80	28.26	

Table 8 (b) Correlation of abnormal palmar creases (left hand) with scores on Indian Scale for Assessment of Autism (ISAA)

Palmar crease left	Cases (n=50)	ISAA so	p value	
hand		Mean	Std. Deviation]
Normal	24	111.08	30.37	0.285
Simian	10	110.30	24.33	Not
Sydney	12	129.42	22.70	signific
Suwon	4	117.00	35.87	ant
	50	115.80	28.26]

Table 2 shows the prevalence of different types of palmar creases in both hands of case and control groups. Majority of children in control group had normal palmar creases. The prevalence of abnormal palmar creases was much higher in the case group than in control group and this difference was statistically significant (p<0.05).

When two groups were compared for abnormal palmar creases in individual hands, the case group had higher prevalence for abnormal creases in both hands as compared to control group and this difference was statistically significant (Table 3). Suwon crease was seen only in left hand of the case group.

The severity of autism spectrum disorder in case group was assessed by Childhood Autism Rating Scale- 2nd edition (CARS). Out of 50 children with autism spectrum disorder, 36 had mild to moderate autism, and 14 had severe autism (Table 5). When patterns of palmar crease of both hands were compared with severity of autism spectrum disorder, the difference was not found to be statistically significant (Table 6).

On assessing severity of autism by Indian Scale for Assessment of Autism (ISAA), out of 50 children with autism spectrum disorder, 15 had mild autism, 27 had moderate and 8 had severe autism (Table 7). When patterns of palmar crease of both hands were compared with severity of autism spectrum disorder, the difference was not found to be statistically significant (Table 8).

However, no correlation of abnormal palmar creases was noticed with gender (Table 4) in both the groups or with severity of the disorder in the case group, as measured by CARS and ISAA (Tables 6-8).

DISCUSSION

The abnormal palmar crease patterns being developmental in nature and that too from neuroectoderm, may point towards the presence of neurodevelopmental disorder such as autism spectrum disorder. Our study was an attempt to estimate the prevalence of different types of abnormal palmar creases in children with autism spectrum disorder and to compare these with those seen in typically developing children. Majority of the children with autism spectrum disorder in our study belonged to the age group of 4-7 years with mean age of 6.5 years. This observation was similar to the findings of other studies. [12-16] In our study, 86% (n=43) children with autism spectrum disorder were males and this is similar to the findings of other studies[17-19] including Indian studies, which also mention the male preponderance in their study samples.[13,20] The finding that majority of children with autism spectrum disorder (64%) in our study belonged to nuclear family (as compared to 58% in control group), is comparable to an Indian study in which 72.4% children with autism spectrum disorder belonged to nuclear family. [21] The overall prevalence of abnormal palmar creases in the case group was 47%, as compared to 14% in control group. Among the abnormal palmar creases seen in the case group, prevalence of Simian crease (22%) was slightly more than the Sydney crease (21%), followed by Suwon crease (4%). Not much of the research has been done on assessing the abnormal palmar creases lately, but one study on abnormal

dermatoglyphics in adolescents with autism spectrum disorders by de Bruin et al reported a prevalence of 2.2% of Sydney crease. [22] We also tried to find correlation of palmar crease patterns in children with autism spectrum disorder to gender and severity of autism but could not find the difference to be statistically significant (p>0.05). To our best of the knowledge no work is done in this area in recent times. We can infer at this stage from these findings that the presence of abnormal palmar creases may point towards the presence of some neurodevelopmental disorder such as autism spectrum disorder but it however, does not signify the severity of autism spectrum disorder.

CONCLUSION

This study is one of its kind in India that assesses the prevalence of different patterns of abnormal palmar creases in children with autism spectrum disorder. The comparison of abnormal palmar crease patterns in children with autism spectrum disorder with that in typically developing children provides further strength to the study results. Although it was a hospital-based study with a small sample size, so we cannot generalize the results into the community. The presence of abnormal palmar creases may prompt clinicians for early screening and early diagnosis and then timely institution of appropriate interventions leading to better outcomes in affected children and alleviation of parental stress and burden

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