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Dermatoglyphics in Down's syndrome

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Abstract

This study aimed at determining the dermatoglyphic patterns of Down's syndrome subjects in Nigeria. A digital scanning method was used to select subjects and a total of 101 Down's syndrome subjects (58 males and 43 females) and 100 control subjects (65 males and 35 females) were used for the study. The subjects were conveniently selected from various special schools in Nigeria. The data were tested using Chi-square and Mann-Whitney U test. A significantly increased ulnar loop was observed for Down's syndrome. The distribution of fingerprint patterns and finger ridge counts were observed to be significantly different between Down's syndrome and control subjects and they were particularly observed in the index and middle fingers (both right and left hands) for both sexes, male subjects (p<0.05). As such, the total finger ridge count was higher in Down's syndrome as compared to control subjects and it was particularly observed on the right hands of both sexes and male subjects (p<0.05). The females showed no significant difference. In conclusion, the study revealed significant difference between Down's syndrome and control subjects deducing that dermatoglyphics could be correlated to Down's syndrome and could be used for its early diagnosis to aid early intervention.

Keywords: Dermatoglyphics, Down's syndrome, Control, Early intervention, Nigeria

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1. Introduction

Dermatoglyphics as the scientific study of epidermal ridges of the skin (both fingerprints and footprints) (Moore and Persaud, 2003), is believed to provide understanding and solution to questions in some areas of life especially in medicine, genetics and evolution (Pratibha *et al.*, 2011; Oladipo *et al.*, 2013). This science of dermatoglyphics is seen to be important because it is based on the fact that ridges are different for different fingers and no two persons show exactly similar fingerprint patterns, also, the ridges are permanent throughout the life of a person (Singh *et al.*, 2016a; Sandeep *et al.*, 2012; Jeewandeep and Arvinder, 2013). Again the development of the brain and the skin at about the same period could be the cause of ridge pattern affected by certain abnormalities of early development. As a result of this effect, dermatoglyphics is correlated with some genetic abnormalities such as mental illnesses and chromosomal disorders such as down syndrome, autism, diabetes, schizophrenia, etc. (Walker, 1977; Lainhart *et al.*, 1997; Bulagouda *et al.*, 2013; Singh *et al.*, 2016a, b).

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Based on these, dermatoglyphics is applied and is used in various fields (Tarca, 2008; Oladipo et al., 2010; Pratibha et al., 2011) such as identification of individual. This application started with William Herschel in 1858 that identified his contractors from their finger prints and hand prints (Cummins and Midlo, 1943). Earlier before now, dermatoglyphics was used to understand individual potentiality(ies) and talents (Campbell, 1998; Sharma et al., 2018). It was believed that everybody inherits some level of innate intelligence from their parents (The Secret of Your Child Fingerprint, 2011). The palmistry uses it for fortune telling as well as prophecy (Campbell, 1998). Today, dermatoglyphics has so much application than what was mentioned above. Its applications can be found in psychiatry, twin diagnosis, forensic and anthropology studies, maternal disputed paternity, as well as medical/disease diagnosis (Schaumann and Alter, 1976). Studies have shown that, dermatoglyphics plays a crucial role in the early diagnosis of genetic diseases (Verbov, 1970; Oladipo et al., 2009; Mollic and Habib, 2011; Reed et al., 1978; Osaat et al., 2022; Osaat et al., 2019)

Even in the phase of DNA testing, fingerprint or dermatoglyphics is still significantly important especially for twin diagnosis. It has been reported that no two monozygotic twins have the same fingerprint pattern and ridges (Reed et al., 1975; Reed et al., 1978). Fingerprint plays its outstanding roles of individuality and uniqueness for proper identification. No wonder dermatoglyphics was referred to as "DNA reflected in the appearance of our body" (The Secret of Your Child's Fingerprint, 2011). Today medical dermatoglyphics applies dermatoglyphics in the diagnosis of so many diseases with 80% to 90% accuracy (Schaumann and Alter, 1976). Infact the diagnosis of these disorders can now be done on the basis of dermatoglyphics analysis alone (Johnny, 2018).

Down's Syndrome (DS) is a chromosomal condition caused by the presence of all or part of a third copy of chromosome twenty one (21). It is also called trisomy 21 (Gordon, 2010; Butler and Meaney, 2005). It is typically associated with a delay in cognition ability (mental retardation) and physical growth with a particular set of facial characteristics. A large number of individuals with Down's syndrome have a severe degree of intellectual disability. The incidence of Down's syndrome in Nigeria as reported by Adeyokunnu (1982) is one in eight hundred and sixty five (865) livebirths in Nigerian hospital.

It is evident that some researchers have reported certain level of correlations between dermatoglyphics and some disorders that have genetic origin such as autism (Milicic *et al.*, 2003; Stosljevic and Adamovic, 2013; Oladipo *et al.*, 2013; Osaat *et al.*, 2019) and Down's syndrome (Boroffice, 1978; Arrieta *et al.*, 1990; Tarca and Barabolski, 2003; Sharma *et al.*, 2012; Stosljevic and Adamovic, 2013), mental retardation (Stevenson *et al.*, 1997), breast cancer (Raizada *et al.*, 2013), Idiopathic Dilated Cardiomyopathy (Oladipo *et al.*, 2007), Obesity (Oladipo *et al.*, 2010), Schizophrenia (Ozyurt *et al.*, 2010), Sickle Cell Anemia (Ramesh *et al.*, 2012), Epilepsy (Aminu *et al.*, 2014), and Diabetes Mellitus (Shield *et al.*, 1995; Oladipo and Ogunnowo, 2004; Oladipo *et al.*, 2012; Pushpa *et al.*, 2013), due to the fact that genetic and uterine environmental events influence dermal pattern formation and so genetic anomalies in the process leave markers in the ridge pattern.

According to Sharma *et al.* (2012) found out that dermatoglyphics may be used as diagnostic tool for predicting the possibilities of development of Down's syndrome at later date. Barbosa *et al.* (2009) also found out that Down's syndrome can be determined using dermatoglyphics.

Boroffice (1978) conducted a study on the dermatoglyphics and found out that Down's syndrome is highly dermatoglyphic specific.

Obviously, dermatoglyphics have been extensively studied in Nigeria and other countries especially on other genetic disorders, however information on dermatoglyphic patterns of Down syndrome are relatively scarce in Nigeria. Therefore, this study aimed at determining the dermatoglyphics (fingerprint) patterns of Down syndrome in Nigeria.

2. Materials and Methods

A descriptive sample survey method was used to investigate the fingerprints patterns of Down's syndrome. The research was carried out in some selected cities in Nigeria such as Lagos, Abuja and Port Harcourt. This study comprised both male and female Down's syndrome subjects in Nigeria, between 5 to 35 years of age. Though no documented statistical record on the population of Down's syndrome subjects in Nigeria, Adeyokunnu (1982) reported the prevalence rate of 1 in 865 live births. Based on this prevalence, the sample

size was determined using Cochran formula (Daniel, 1999) and the minimum sample size of Down's syndrome was 49, however the sample size used for this research study was 101 (58 males and 43 females) for Down's syndrome. The subjects who volunteered through their parents or institutional authorities to participate in the study, with no form of trauma or anomaly in their palms and feet and must be within the age range for the study were selected for the study.

The study adopted the method according to Oghenemavwe and Osaat (2015). The dermatoglyphic patterns were collected and determined using the scanning method which involves a High-resolution digital scanner-G3110 Scanjet Scanner with 4800x9600 dpi resolution connected to a laptop to identify and classify dermatoglyphics. The scanner and laptop were both electrically powered using any electrical source.

The subjects' fingers and palms were thoroughly washed with water and soap and dried with clean towel to remove dirt. The subject was asked or assisted to place the washed palms on the scanner and accordingly the palms were scanned. The thumb prints were taken separately from the other fingers and palm because of its position to obtain maximum clarity. The scanned images were saved in a folder and named appropriately. Later on, collation of raw data was obtained from the scan images and used for further analysis. Ridge counting was done using AUTOCAD Program (version 2010), it was also used to count ridges with limited errors. The Finger Print Patterns analyzed include: Arch (A), Ulnar Loop (UL), Radial Loop (RL) and Whorl (W). Finger Ridge Count (FRC) and Total Finger Ridge Count (TFRC) were analyzed for individual fingers on both right and left hands. TFRC includes the sum of the ten finger ridge counts. This counting was done along the straight line connecting one tri-radial point (tri-radial point is formed by the confluence of the three ridge system) to the point of core.

The data obtained from this study were subjected to test using Statistical Package for Social Science ((SPSS) IBM @ Version 23 New York). For clarity, tables were used to present results. Mann-Whitney U test analysis was used to compare all the quantitative data such as Finger Ridge Counts (FRC), while Chi-square was used for analysis on percentage frequencies of finger patterns. All statistical testing was done at 95% confidence level with p-value less than 0.05 (p<0.05) taken to be significant.

Prior to commencement of the research work, ethical approval was sought from the Research Ethics Committee of the School of Graduate Studies, University of Port Harcourt in form of proposal writing and it was approved with reference number UPH/CEREMAD/REC/04. In addition, informed consent was obtained from the parents/guidance and institutional authorities of the subjects by signing a consent form given to them before samples of the subjects under study were taken.

3. Results

Tables 1a and 1b showed the distribution of right and left finger print patterns and test of association respectively, in both sexes of Down's syndrome and normal subjects. A significant difference in the index, middle and little fingers of the right hand of Down's syndrome and normal subjects (p<0.05) was observed. The thumb and ring fingers were not statistically significant though they have variable percentage of patterns in those fingers. Table 1c summarized the differences in the pattern of Down's syndrome and normal subjects as tested using chi square. In both right and left hands there were significant differences amongst the patterns of Down's syndrome and normal subjects (p<0.05).

In Tables 2a and 2b, significant difference was observed in males of Down's syndrome and normal subjects in both the index, middle and little fingers of the right and left hands (p<0.05). Other fingers were not significant as p>0.05. In Table 2c, the result showed that there was statistically significant difference in the fingerprint patterns of Down's syndrome and normal male subjects on both right and left hands (p<0.05).

However, in Table 3a the female showed significant differences only in the middle fingers on the right hand of Down's syndrome and normal subjects (p<0.05). The other fingers were not statistically significant though they were variable percentage of patterns in those fingers. In Table 3b, on the left hand, significant difference was demonstrated in the index and middle fingers of Down's syndrome and normal subjects (p<0.05). The thumb, ring and little fingers were not statistically significant as well. Table 3c summarized the differences in the pattern of Down's syndrome and normal subject as tested using chi square. Like the male subjects, there

Table 1a: Dis	stribution of th	e right fing	ger print pat	tern and tes	t of associati	on of both	sexes		
Dialet Geran	Canada	A 1- (0/)	Radial	Ulnar	TA71s out (0/)	Chi-s	square analysis		
Right finger	Group	Arch (%)	loop (%)	loop (%)	Whorl (%)	X ²	df	P-value	
Thumb	DS subjects	16 (15.8)	-	47 (46.5)	38 (37.6)	4.20	2	0.12	
Inumb	NO subjects	12 (12.0)	-	36 (36.0)	52 (52.0)	4.20		0.12	
In day.	DS subjects	6 (5.9)	1 (1.0)	84 (83.2)	10 (9.9)	40.2F	3	0.00**	
Index	NO subjects	15 (15.0)	2 (2.0)	40 (40.0)	43 (43.0)	40.35	3	0.00***	
Middle	DS subjects	-	-	93 (92.1)	8 (7.9)	18.72	3	0.00**	
Middle	NO subjects	10 (10.0)	1 (1.0)	70 (70.0)	19 (19.0)	16.72		0.00	
Diag	DS subjects	1 (1.0)	-	55 (54.5)	45 (44.6)	1 20	3	0.71	
Ring	NO subjects	2 (2.0)	1 (1.0)	54 (54.0)	43 (43.0)	1.38	3	0.71	
T :ul -	DS subjects	-	-	78 (77.2)	23 (22.8)	12.06	2	0.00**	
Little	NO subjects	3 (3.0)	-	91 (91.0)	6 (6.0)	13.96	2	0.00**	
Note: DS - Down's syndrome; NO - Normal; df - degree of freedom.									

I of t 6:	Canada	A 10 (0/)	Radial	Ulnar	1471a a = 1 (0/)	Chi-	nalysis		
Left finger	Group	Arch (%)	loop (%)	loop (%)	Whorl (%)	X ²	df	P-value	
Thurse	DS subjects	26 (25.7)	-	44 (43.6)	31 (30.7)	5.25	2	0.07	
Thumb	NO subjects	14 (14.0)	-	44 (44.0)	42 (42.0)	5.25	2	0.07	
T., 1.,	DS subjects	2 (2.0)	1 (1.0)	83 (82.2)	15 (14.9)	20.66	3	0.00**	
Index	NO subjects	16 (16.0)	4 (4.0)	46 (46.0)	34 (34.0)	30.66	3	0.00	
Middle	DS subjects	2 (2.0)	-	90 (89.1)	9 (8.9)	16.45	3	0.00**	
Middle	NO subjects	16 (16.0)	1 (1.0)	68 (68.0)	15 (15.0)	16.45	3	0.00***	
D:	DS subjects	2 (2.0)	2 (2.0)	57 (56.4)	40 (39.6)	0.54	2	0.01	
Ring	NO subjects	3 (3.0)	1 (1.0)	57 (57.0)	39 (39.0)	0.54	3	0.91	
T tod.	DS subjects	-	-	84 (83.2)	17 (16.8)	11 50	2	0.00**	
Little	NO subjects	3 (3.0)	-	93 (93.0)	4 (4.0)	11.50	2	0.00**	

Table 1c: Chi-square test comparing the dermatoglyphic patterns of Down's syndrome and normal subjects of both sexes															
C]	Right		Chi-square analysis				Left				Chi-square analysis		
Group	A	RL	UL	W	df	df X ² P-value			RL	UL	W	df	X2	P-value	
DS subjects	23	1	357	124	2	14.17	0.00**	32	3	358	112	3	11.46	0.01**	
NO subjects 42 4 302 149 3	3	14.17	0.00**	52	6	308	134	3	11.40	0.01					
Notes A Augle	Note: A Auch DI Padial Ican III Illner Ican W Wheel ** significant DC Davin's grandrome: NO Normal of														

Note: A - Arch; RL - Radial loop; UL - Ulnar loop; W - Whorl; ** - significant; DS - Down's syndrome; NO - Normal; df - degree of freedom.

were significant differences amongst the patterns of Down's syndrome and normal subjects (p<0.05) in both right and left hands of female subjects.

As shown in Tables 4a and 4b, Mann-Whitney U test was used to test for differences between finger ridge count of Down's syndrome subjects and normal subjects on the right and left hands respectively of both sexes.

Table 2a: Distribution of the right finger print pattern and test of association in males of Down's	
syndrome and normal subjects	

Dight fings	Croun	A == (0/)	Radial loop	Ulnar loop	147h out (0/)	Chi-sq	uare aı	nalysis
Right finger	Group	Arch (%)	(%)	(%)	Whorl (%)	X^2	df	P-value
Thumb	DS subjects	11 (19.0)		25 (43.1)	22 (37.9)	4.00	2	0.13
Thumb	Normal subjects	7 (10.8)		22 (33.8)	36 (55.4)	4.08		0.13
Index	DS subjects	6 (10.3)		45 (77.6)	7 (12.1)	22.14	3	0.00**
maex	Normal subjects	11 (16.9)	2 (3.1)	2 (3.1) 24 (36.9)		22.14	5	0.00
Middle	DS subjects			52 (89.7)	6 (10.3)	12.50	2	0.01**
Middle	Normal subjects	9 (13.8)	1 (1.5)	43 (66.2)	12 (18.5)	12.50	3	0.01
Dina	DS subjects			31 (53.4)	27 (46.6)	3.22	3	0.36
Ring	Normal subjects	2 (3.1)	1 (1.5)	37 (56.9)	25 (38.5)	3.22	3	0.36
T :ul -	DS subjects			41 (70.0)	17 (29.3)	12 (12	2	0.00**
Little	Little Normal subjects			58 (89.2)	4 (6.2)	13.613	2	0.00**
Note: DS - Down's syndrome; NO - Normal; df - degree of freedom.								

Table 2b: Distribution of t	he left finger print pattern and test of association in males of Down's
syndrome and normal sub	jects

Group	Group	Group	Group	Group	Group	Group Arch (%)		Radial loop Ulnar loop		TA71- a-1 (0/)	Chi-square analysis		
Gloup	Arch (%)	(%)	(%)	Whorl (%)	X ²	df	P-value						
DS subjects	17 (29.3)		23 (39.7)	18 (31.0)	E EO	2	0.06						
Normal subjects	8 (12.3)		30 (46.2)			2	0.06						
DS subjects 1			49 (84.5)	8 (13.8)	24.94	2	0.00**						
Normal subjects	13 (20.0)	4 (6.2)	28 (43.1)	20 (30.8)	24.84	3	0.00						
DS subjects	2 (3.4)		51 (87.9)	5 (8.6)	0.13	2	0.04**						
Normal subjects	12 (18.5)	1 (1.5)	46 (70.8)	6 (9.2)	0.12		0.04						
DS subjects			33 (56.9)	25 (43.1)	E E2	2	0.14						
Normal subjects	3 (4.6)	1 (1.5)	42 (64.6)	19 (29.2)	3.32	3	0.14						
DS subjects			47 (81.0)	11 (19.0)	9.56	2	0.01**						
Normal subjects	3 (4.6)		59 (90.8)	3 (4.6)	0.36	2	0.01***						
1	formal subjects DS subjects OS subjects	formal subjects 8 (12.3) DS subjects 1 (1.7) formal subjects 13 (20.0) DS subjects 2 (3.4) formal subjects 12 (18.5) DS subjects formal subjects 3 (4.6) DS subjects formal subjects 3 (4.6)	formal subjects 8 (12.3) DS subjects 1 (1.7) formal subjects 13 (20.0) 4 (6.2) DS subjects 2 (3.4) formal subjects 12 (18.5) 1 (1.5) DS subjects formal subjects 3 (4.6) 1 (1.5) DS subjects formal subjects 3 (4.6)	Jormal subjects 8 (12.3) 30 (46.2) DS subjects 1 (1.7) 49 (84.5) Jormal subjects 13 (20.0) 4 (6.2) 28 (43.1) DS subjects 2 (3.4) 51 (87.9) Jormal subjects 12 (18.5) 1 (1.5) 46 (70.8) DS subjects 33 (56.9) Jormal subjects 3 (4.6) 1 (1.5) 42 (64.6) DS subjects 47 (81.0)	formal subjects 8 (12.3) 30 (46.2) 27 (41.5) DS subjects 1 (1.7) 49 (84.5) 8 (13.8) formal subjects 13 (20.0) 4 (6.2) 28 (43.1) 20 (30.8) DS subjects 2 (3.4) 51 (87.9) 5 (8.6) formal subjects 12 (18.5) 1 (1.5) 46 (70.8) 6 (9.2) DS subjects 33 (56.9) 25 (43.1) formal subjects 3 (4.6) 1 (1.5) 42 (64.6) 19 (29.2) DS subjects 47 (81.0) 11 (19.0) 10 (19.0) formal subjects 3 (4.6) 59 (90.8) 3 (4.6)	5.58 5.58	Jornal subjects 8 (12.3) 30 (46.2) 27 (41.5) 27 (41.5) DS subjects 1 (1.7) 49 (84.5) 8 (13.8) 24.84 3 Jornal subjects 13 (20.0) 4 (6.2) 28 (43.1) 20 (30.8) 24.84 3 DS subjects 2 (3.4) 51 (87.9) 5 (8.6) 8.12 3 Jornal subjects 12 (18.5) 1 (1.5) 46 (70.8) 6 (9.2) 8.12 3 DS subjects 3 (4.6) 1 (1.5) 42 (64.6) 19 (29.2) 5.52 3 DS subjects 47 (81.0) 11 (19.0) 8.56 2 Jornal subjects 3 (4.6) 59 (90.8) 3 (4.6) 8.56 2						

 $\mbox{\bf Note:}\ \mbox{DS}$ - Down's syndrome; NO - Normal; df - degree of freedom.

Table 2c: Chi-square test comparing the finger patterns of male Down's syndrome and normal subjects

Group		Ri	ght		Chi-square analysis		Left				Chi-square analysis			
Group	A	RL	UL	W	df	X2	P-value	A	RL	UL	W	df	X2	P-value
DS subjects	17	4	194	79	2	11.37	0.010**	20	-	203	67	2	10.62	0.014**
NO subjects	151	-	194	79	3	11.57	0.010	39	6	205	75	3	10.62	0.014

Note: A - Arch; RL - Radial loop; UL - Ulnar loop; W - Whorl; ** - significant; DS - Down's syndrome; NO - Normal; df - degree of freedom.

In Table 4a, the result on the right hand showed that the index, middle and little fingers of Down's syndrome subjects were significantly increased from those of normal subjects (p<0.05), while in Table 4b the result on the

Table 3a: Distribution of the right finger print patterns and test of association in females of Down's syndrome and normal subjects

D:-1-1 C:	C	A1- (0/)	Radial	Ulnar loop	TATI ::1 (0/)	Chi-se	quare an	alysis
Right finger	Group	Arch (%)	loop (%)	(%)	Whorl (%)	X^2	df	P-value
Therenale	DS subjects	5 (11.6)		22 (51.2)	16 (37.2)	0.07	2	0.62
Thumb	NO subjects	5 (14.3)		14 (40.0)	16 (45.7)	0.97	2	0.62
Index	DS Subjects		1 (2.3)	39 (90.7)	3 (7.0)	22.03	3	0.00**
muex	NO subjects	4 (11.4)		16 (45.7)	15 (42.9)	22.03	3	0.00
Middle	DS Subjects			41 (95.3)	2 (4.7)	F 00	2	0.05
Middle	NO subjects	1 (2.9)		27 (77.1)	7 (20.0)	5.90	2	0.05
Dia a	DS subjects	1 (2.3)		24 (55.8)	18 (41.9)	1.39	2	0.50
Ring	NO subjects			17 (48.6)	18 (51.4)	1.39	2	0.50
T :u1.	DS subjects			37 (86.0)	6 (14.0)	1 40	1	0.20
Little	NO subjects			33 (94.3)	2 (5.7)	1.42	1	0.28
Note: DS - Dow	vn's syndrome;	NO - Normal;	df - degree o	of freedom.				•

Table 3b: Distribution of the left finger print patterns and test of association in females of Down's syndrome and normal subjects

I - 6: 6:	C	A1- (0/)	Radial	Ulnar loop	TA71 ::1 (0/)	Chi-sq	quare analysis	
Left finger	Group	Arch (%)	loop (%)	(%)	Whorl (%)	X^2	df	P-value
Thumb	DS subjects	9 (20.9)		21 (48.8)	13 (30.2)	1.34	2	0.51
Thumb	NO subjects	6 (17.1)		14 (40.0)	15 (42.9)	1.34	2	0.51
Index	DS subjects	1 (2.3)	1 (2.3)	34 (79.1)	7 (16.3)	8.53	3	0.04**
muex	NO subjects	3 (8.6)		18 (51.4)	14 (40.0)	6.33	3	0.04
Middle	DS subjects			39 (90.7)	4 (9.3)	9.95	2	0.01**
Middle	NO subjects	4 (11.4)		22 (62.9)	9 (25.7)	9.93		0.01
Ring	DS subjects	2 (4.7)	2 (4.7)	24 (55.8)	15 (34.9)	6.03	3	0.11
King	NO subjects			15 (42.9)	20 (57.1)	6.03	3	0.11
Little	DS subjects			37 (86.0)	6 (14.0)	2.91	1	0.12
Little	NO subjects			34 (97.1)	1 (2.9)	2.91	1	0.12
N. A. D.C. D.		NO NI 1	16 1	1	·		·	

Note: DS - Down's syndrome; NO - Normal; df - degree of freedom.

Table 3c: Chi-square test comparing the finger patterns of female Down's syndrome and normal subjects

Сиона		Ri	ght		Chi	Chi-square analysis			Left				Chi-square analysis		
Group	A	RL	UL	W	df	X ²	P-value	A	RL	UL	W	df	X2	P-value	
DS subjects	6	1	163	45	2	11.27	7 0.010**	12	3	155	45	2	11.42	0.010**	
NO subjects	10	-	107	58	3	11.2/	0.010	13	-	103	59)	11.42	0.010	

Note: A - Arch; RL - Radial loop; UL - Ulnar loop; W - Whorl; ** - significant; DS - Down's syndrome; NO - Normal; df - degree of freedom.

left hand showed only the index and middle fingers were significant (p<0.05). In Tables 5a and 5b, the results on the right and left hands respectively of male Down's syndrome and normal subjects showed that the index

Table 4a: Mann-Whitney U test comparing the right finger ridge count of Down's syndrome and
normal Subjects of both sexes

Right finger ridge count	Group	N	Mean rank	Sum of ranks	Mann- Whitney U	Wilcoxon W	Z	P-value
Thumb	DS subjects	101	103.77	10481.00	4770.00	9820.00	-0.68	0.50
Tnumb	NO subjects	100	98.20	9820.00	4//0.00	9820.00	-0.68	0.50
Indov	DS subjects	101	117.78	11896.00	3355.00	8405.00	-4.12	0.00**
Index	NO subjects	100	84.05	8405.00	3333.00	0405.00	-4 .12	0.00
Middle	DS subjects	101	122.33	12355.50	- 2895.50	7945.50	-5.24	0.00**
Middle	NO subjects	100	79.46	7945.50	2095.50	7945.50		
Dina	DS subjects	101	103.34	10437.50	4813.50	9863.50	-0.57	0.57
Ring	NO subjects	100	98.64	9863.50	4013.30	9003.30	-0.57	0.57
T :ul.	DS subjects	101	109.65	11075.00	44.774.00	0226.00	0.10	0.02**
Little	NO subjects	100	92.26	9226.00	4176.00	9226.00	-2.13	0.03**

Note: ** - Significant; DS - Down's syndrome; NO - Normal; z - z score.

Table 4b: Mann-Whitney U test comparing the left finger ridge count of Down's syndrome and normal Subjects of both sexes

Group	N	Mean rank	Sum of ranks	Mann- Whitney U	Wilcoxon W	Z	P-value
DS subjects	101	100.74	10175.00	E024.00	10175.00	-0.06	2.05
NO subjects	100	101.26	10126.00	5024.00	10175.00		0.95
DS subjects	101	116.74	11791.00	2460.00	9E10.00	2.07	0.00**
NO subjects	100	85.10	8510.00	3460.00	8510.00	-3.67	0.00
DS subjects 10	101	118.23	11941.00	2210.00	9260.00	-4.23	0.00**
NO subjects	100	83.60	8360.00	3310.00	6360.00		
DS subjects	101	95.73	9668.50	4F17 F0	0668.50	1.20	0.20
NO subjects	100	106.33	10632.50	4517.50	9668.30	-1.29	0.20
DS subjects	101	100.40	10140.50	4000 50	10140 50	0.15	0.88
NO subjects	100	101.61	10160.50	4989.30	10140.50	-0.15	
	DS subjects NO subjects DS subjects NO subjects DS subjects NO subjects NO subjects DS subjects DS subjects Solve the subjects Solve the subjects DS subjects Solve the subjects	DS subjects 101 NO subjects 100 DS subjects 100 NO subjects 100 DS subjects 101 NO subjects 100 DS subjects 100 DS subjects 101 NO subjects 101 NO subjects 100 DS subjects 100 DS subjects 100	Group N rank DS subjects 101 100.74 NO subjects 100 101.26 DS subjects 101 116.74 NO subjects 100 85.10 DS subjects 101 118.23 NO subjects 100 83.60 DS subjects 101 95.73 NO subjects 100 106.33 DS subjects 101 100.40	Group N rank ranks DS subjects 101 100.74 10175.00 NO subjects 100 101.26 10126.00 DS subjects 101 116.74 11791.00 NO subjects 100 85.10 8510.00 DS subjects 101 118.23 11941.00 NO subjects 100 83.60 8360.00 DS subjects 101 95.73 9668.50 NO subjects 100 106.33 10632.50 DS subjects 101 100.40 10140.50	Group N rank ranks Whitney U DS subjects 101 100.74 10175.00 5024.00 NO subjects 100 101.26 10126.00 3024.00 DS subjects 101 116.74 11791.00 3460.00 NO subjects 100 85.10 8510.00 3310.00 NO subjects 100 83.60 8360.00 3310.00 DS subjects 101 95.73 9668.50 4517.50 NO subjects 100 106.33 10632.50 4517.50 DS subjects 101 100.40 10140.50 4989.50	Group N rank ranks Whitney U W DS subjects 101 100.74 10175.00 5024.00 10175.00 NO subjects 100 101.26 10126.00 324.00 10175.00 DS subjects 101 116.74 11791.00 3460.00 8510.00 NO subjects 100 85.10 8510.00 3310.00 8360.00 NO subjects 100 83.60 8360.00 3310.00 8360.00 DS subjects 101 95.73 9668.50 4517.50 9668.50 NO subjects 100 106.33 10632.50 4517.50 9668.50 DS subjects 101 100.40 10140.50 4989.50 10140.50	Group N rank ranks Whitney U W Z DS subjects 101 100.74 10175.00 5024.00 10175.00 -0.06 NO subjects 100 101.26 10126.00 3024.00 10175.00 -0.06 DS subjects 101 116.74 11791.00 3460.00 8510.00 -3.87 NO subjects 100 85.10 8510.00 3310.00 8360.00 -4.23 NO subjects 101 95.73 9668.50 4517.50 9668.50 -1.29 NO subjects 101 106.33 10632.50 4517.50 9668.50 -1.29 DS subjects 101 100.40 10140.50 4989.50 10140.50 -0.15

Note: ** - Significant; DS - Down's syndrome; NO - Normal; z - z score.

Table 4c: Total Finger Ridge Count (TFRC) of Down's syndrome and normal subjects compared using Mann-Whitney U test of both sexes

Total finger ridge count	Group	N	Mean rank	Sum of ranks	Mann- Whitney U	Wilcoxon W	Z	P-value
Right total RC	DS subjects	101	112.45	11357.00	3894.00	8944.00	-2.80	0.01**
Right total KC	NO subjects	100	89.44	8944.00	3694.00			
I () 1 DC	DS subjects	101	107.33	10840.50	4410.50	0460.50	1 55	0.10
Left total RC	NO subjects	100	94.61	9460.50	4410.30	9460.50	-1.55	0.12

Note: ** - Significant; DS - Down's syndrome; RC - Ridge count; z - z score.

and middle fingers were significant different in their ridge counts (p<0.05). In Table 6a only the little finger was significantly different in their ridge count between female Down's syndrome and female normal subjects

Table 5a: Distribution of the right finger ridge count and test of association in males of Down's
syndrome and normal subjects

Right finger	Group	N	Mean	Sum of	Mann-	Wilcoxon	Z	P-value
ridge count	r	-,	rank	ranks	Whitney U	W	_	
Thumb	DS subjects	101	103.77	10481.00	4770.00	9820.00	-0.68	0.50
THUIHD	NO subjects	100	98.20	9820.00	4770.00	9620.00	-0.00	0.50
Index	DS subjects	101	117.78	11896.00	3355.00	8405.00	-4.12	0.00**
index	NO subjects	100	84.05	8405.00	3333.00	0405.00		0.00
Middle	DS subjects	101	122.33	12355.50	2895.50	7945.50	-5.24	0.00**
Middle	NO subjects	100	79.46	7945.50	2095.50	7945.50		
Dina	DS subjects	101	103.34	10437.50	4813.50	9863.50	-0.57	0.57
Ring	NO subjects	100	98.64	9863.50	4015.50	9665.50	-0.37	0.57
T:ul-	DS subjects	101	109.65	11075.00	4177,00	0226.00	2.12	0.03**
Little	NO subjects	100	92.26	9226.00	4176.00	9226.00	-2.13	

Note: ** - Significant; DS - Down's syndrome; NO - Normal; z - z score.

Table 5b: Distribution of the left finger ridge count and test of association in males of Down's syndrome and normal subjects

Right finger ridge count	Group	N	Mean rank	Sum of ranks	Mann- Whitney U	Wilcoxon W	z	P-value
Thumb	DS finger	58	64.37	3733.50	1747.50	2002 50	0.70	0.40
	NO finger	65	59.88	3892.50	1747.50	3892.50	-0.70	0.48
Index	DS finger	58	75.08	4354.50	1126.50	3271.50	2.07	0.00**
index	NO finger	65	50.33	3271.50	1120.50	32/1.50	-3.86	0.00
Middle	DS finger	58	78.80	4570.50	010.50	3055.50	-4.95	0.00**
Ivildale	NO finger	65	47.01	3055.50	910.50	3033.30	-4.93	
Dia -	DS finger	58	66.02	3829.00	1(52.00	3797.00	1 10	0.24
Ring	NO finger	65	58.42	3797.00	1652.00	3/9/.00	-1.18	0.24
Little	DS finger	58	65.28	3786.00	1695.00	3840.00	-0.97	0.22
Little	NO finger	65	59.08	3840.00	1093.00	3640.00	-0.97	0.33

Note: ** - Significant; DS - Down's syndrome; NO - Normal; z - z score.

Table 5c: Male Total Finger Ridge Count (TFRC) of Down's syndrome and Normal subjects compared using Mann-Whitney U test

Group	N	Mean rank	Sum of ranks	Mann- Whitney U	Wilcoxon W	Z	P-value
DS subjects	58	71.14	4126.00	1355.00	2500.00	2 (0	0.01**
NO subjects	65	53.85	3500.00	1555.00	5500.00	-2.09	
DS subjects	58	66.06	3831.50	1640 50	2704 50	1 10	0.23
NO subjects 65 58.38 3794.50	1049.50	3794.30	-1.19	0.23			
	DS subjects NO subjects DS subjects	DS subjects 58 NO subjects 65 DS subjects 58	DS subjects 58 71.14 NO subjects 65 53.85 DS subjects 58 66.06	Group N rank ranks DS subjects 58 71.14 4126.00 NO subjects 65 53.85 3500.00 DS subjects 58 66.06 3831.50	Group N rank ranks Whitney U DS subjects 58 71.14 4126.00 NO subjects 65 53.85 3500.00 DS subjects 58 66.06 3831.50 1649.50	Group N rank ranks Whitney U Wilcoxon W DS subjects 58 71.14 4126.00 1355.00 3500.00 NO subjects 65 53.85 3500.00 3500.00 3500.00 DS subjects 58 66.06 3831.50 1649.50 3794.50	Group N rank ranks Whitney U Wilcoxon W Z DS subjects 58 71.14 4126.00 1355.00 3500.00 -2.69 NO subjects 65 53.85 3500.00 1649.50 3794.50 -1.19

Note: ** - Significant; DS - Down's syndrome; RC - Ridge count; z - z score.

on the right, while in Table 6b, on the left hand the middle finger of female Down's syndrome and normal subjects showed significant difference in their ridge counts (p<0.05).

Table 6a: Distribution of the right I	inger Ridge Count (FRC) and test of association in females of
Down's syndrome and normal sub	jects

Right finger ridge count	Group	N	Mean rank	Sum of ranks	Mann- Whitney U	Wilcoxon W	Z	P-value
Thumb ———	DS finger	43	40.87	1757.50	(02 F0	1222 FO	0.50	
	NO finger	35	37.81	1323.50	693.50	1323.50	-0.59	0.55
Index	DS finger	43	44.00	1892.00	559.00	1189.00	-1.95	0.05
inaex	NO finger		1189.00	-1.93	0.05			
Middle	DS finger	43	43.74	1881.00	F70 00	1200.00	-1.85	0.06
Middle	NO finger	35	34.29	1200.00	570.00	1200.00	-1.65	0.06
D:	DS finger	43	38.05	1636.00	(00.00	1636.00	0.62	0.53
Ring	NO finger	35	41.29	1445.00	690.00	1636.00	-0.63	0.55
I :ulo	DS finger	43	45.28	1947.00	504.00	1134.00	2.50	0.01**
Little	NO finger	35	32.40	1134.00		1134.00	-2.50	0.01***

Note: ** - Significant; DS - Down's syndrome; NO - Normal; z - z score.

Table 6b: Distribution of the left Finger Ridge Count (FRC) and test of association in females of Down's syndrome and normal subjects

Left finger ridge count	Group	N	Mean rank	Sum of ranks	Mann- Whitney U	Wilcoxon W	Z	P-value
TI 1	DS finger	43	38.02	1635.00	600.00	1635.00	-0.64	
Thumb	NO finger	35	41.31	1446.00	689.00	1655.00		0.52
Index	DS finger	43	43.87	1886.50	F(4 F0	1194.50	-1.89	0.06
index	NO finger	35	34.13	1194.50	564.50			0.06
Middle	DS finger	43	45.36	1950.50	F00 F0	1120 50	2.54	0.01**
Middle	NO finger	35	32.30	1130.50	500.50	1130.50	-2.54	0.01***
Dina	DS finger	43	39.60	1703.00	749.00	1279.00	0.05	0.00
Ring	NO finger	35	39.37	1378.00	748.00	1378.00	-0.05	0.96
Little	DS finger	43	41.52	1785.50	665.50	1295.50	0.00	0.28
Little	NO finger	35	37.01	1295.50	003.30	1293.30	-0.88	0.38

Note: ** - Significant; DS - Down's syndrome; NO - Normal; z - z score.

Table 6c: Female Total Finger Ridge Count (TFRC) of Down's syndrome and Normal subjects compared using Mann-Whitney U test

Total finger ridge count	Group	N	Mean rank	Sum of ranks	Mann- Whitney U	Wilcoxon W	Z	P-value
Dialettatal DC	DS subjects	43	42.17	1813.50	(27.50	1267.50	116	0.25
Right total RC	NO subjects	35	36.21	1267.50	637.50		-1.16	0.23
Loft total BC	DS subjects	43	41.78	1796.50	654.50	1284.50	-0.98	0.32
Left total RC	NO subjects	35	36.70	1284.50	054.50	1204.50	-0.96	0.32

Note: ** - Significant; DS - Down's syndrome; RC - Ridge count; z - z score.

In Table 4c, the TFRC result revealed a higher TFRC for Down's syndrome than normal subjects on the both hands though significant on the right hand. In Table 5c, the male Down's syndrome subjects have significantly

higher TFRC on the right hand than the male normal subjects (p<0.05). In Table 6c the female Down's syndrome subjects and normal subjects showed no statistically significant different on both hands though Down's syndrome has a higher TFRC than normal subjects (p>0.05).

4. Discussion

From the result of the study, Down's syndrome had higher percentage of ulnar loop but least percentages of arches and whorls than control subjects on both hands for both sexes, a dermatoglyphic feature which means that, low count of arches and higher count of ulnar loop is associated with Down's syndrome. It suggests genetics in the etiology of Down's syndrome. According to Babler (1991) the type of ridge pattern formed is associated with the timing of primary ridge formation. While early ridge formation was correlated with a whorl pattern, intermediate ridge formation was correlated with a loop and late ridge formation was correlated with an arch pattern. The female subjects have more arches and lesser whorls than their male counterparts. This may suggest sexual dimorphism. Verbov (1970) reported the same finding.

Down's syndrome subjects have decreased percentage of radial loop and it was seen mostly on the right index and left ring fingers. Bryant *et al.* (1970), Plato *et al.* (1973) and Fogle (1990) have similar findings. The frequency of radial loop was more in female than the male counterparts. This suggests that females may be more influenced by genetic factors than environmental factors.

Down's syndrome recorded the least percentage frequency of Whorl pattern which was in line with the study of Bryant *et al.* (1970). This result suggests the association of digital whorl pattern to normal subjects. As said earlier, whorl pattern was associated with early ridge formation (Babler, 1991) and suggests the least influenced by genetic factors. However, a genetic theory postulated by Slatis *et al.* (1976) proposed a genetic theory that ulnar loop is the basic fingerprint pattern found in all the fingers and that other patterns are formed with the activity of various genes acting on the pattern sequence causing deviation from the original pattern.

Again the present study observed an increased percentage of ulnar loops on the hands of Down's syndrome subjects. Infact Down's syndrome was seen to have the highest percentage of ulnar loop than the control. This finding was in line with previous authors (Bryant *et al.*, 1970; Boroffice, 1978; Rajangam *et al.*, 1995) who reported high frequency of ulnar loop on the fingers of Down's syndrome. Ulnar loop was seen on all fingers particularly little finger bilaterally (Bryant *et al.*, 1970; Tarca and Barabolski, 2003). The male Down's syndrome subjects have higher percentage of ulnar loop than female subjects (Barbosa *et al.*, 2009). A study on mentally retarded subjects also recorded strikingly high percentage of ulnar loop as compared with controls (Kiran *et al.*, 2010). The close resemblance could be because individual with Down's syndrome expressed very high level of mental retardation (Villar and Epstein, 2005). This again could serve as a diagnostic/screening tool for Down's syndrome and mentally retarded individuals meaning that though the ulnar loop pattern is the basic fingerprint (Slatis *et al.*, 1976) or the intermediate ridge formation (Babler, 1991), the extreme of it could be abnormal as seen in both Down syndrome subjects and mentally retarded individuals. The predominance of ulnar loop on the hands of Down's syndrome as compared with the control could serve as a dermatoglyphic features for its diagnosis.

The distribution of finger patterns between Down's syndrome and normal subjects on both sexes was significant in index, middle and little fingers for both hands p<0.05. The distribution of finger pattern between the male Down's syndrome and male control subjects was significant in the index, middle and little fingers on both hands, while the female subjects have significant distribution on the index finger alone (right hand) and index and middle fingers (left hand). Summarily, chi-square test showed statistically significant difference between Down's syndrome and control on both right and left finger patterns distribution on both sexes, male and female subjects. It can be deduced from the present study that amongst all fingers, the index finger of Down syndrome subjects is of great importance as it could give insight to finger that deviation from normal is mostly seen, seeing that, the distribution of pattern in index finger is significant for both sexes, for male, and for females subjects as against controls on both hands as seen in the result.

The genetics of finger ridge count and total ridge counts have revealed that there are considerable variations among individuals who are not related, however, statistically positive correlation is observed among relatives (Fogle, 1990). This is as a result of the degree of shared genetic heritage. Traits like TRC have a range of

phenotypic expression and are called quantitative traits. They are capable of letting known the genotype of an individual (Fogle, 1990). Ridge count distribution is a function of the frequency of the pattern types that occur on each of the ten fingers (Holt, 1968). From the present study the finger ridge count of Down's syndrome and controls of both sexes was observed to be significant in the index, middle and little fingers on the right hand. On the left hand only the index and middle fingers were statistically significantly different from each other. This appears to result from the high percentage frequencies of ulnar loop seen in the index, middle and little fingers of Down's syndrome when compared to the normal subjects. As such the Total Finger Ridge Count (TFRC) on the right hand was significantly increased in Down's syndrome when compared with controls while the left hand, statistically insignificant result was observed. This result is in line with Malla and Srivastava (2008) who reported a high TRC for Down's syndrome than control subjects. For the male Down's syndrome subjects, the difference was observed in the index and middle fingers bilaterally and the TFRC was only significant on the right in favour of Down's syndrome. For the female subjects, finger ridge count was significant in the right little finger and the left middle finger. From the result it is obvious that the index finger is the most variable digit and little finger the least due to the large proportion of loop in the little finger. Holt 1968, observed same findings. Despite the differences observed in the female ridge fingers, the TFRC of the female Down's syndrome and normal was statistically insignificant on both right and left hands. This is because the female Down's syndrome subjects have more of arches on their fingers and lower whorls than female controls and the male subjects as well leading to low insignificant TFRC when compared with controls or male subjects (Holt, 1968; Verbov, 1970). The variation in the TFRC may be the result of genetic influence on Down's syndrome.

Figure 1 was the schematic diagram showing the summary of the dermatoglyphics features on the palm for both Down's syndrome and control subjects.

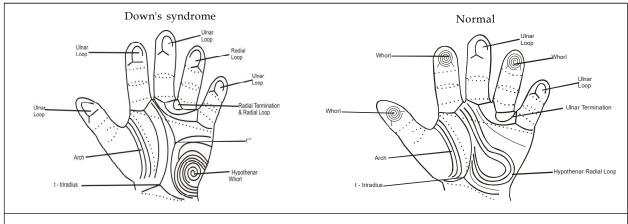


Figure 1: Diagnostic map for Down's syndrome and normal subjects

5. Conclusion

In conclusion, Down's syndrome as compared with the control subjects had an increased ulnar loop on all fingers except finger IV, higher radial loop on the ring finger. Finger ridge pattern and finger ridge count were observed to be significant in the index and middle fingers. These two fingers also are the most variable fingers in the study. High TFRC was observed in both sexes and males subjects except for the female subjects because of the high arches seen on their fingers and these differences are significant on the right hand.

From the results, a strong correlation was observed between dermatoglyphics of Down's syndrome and controls on almost all the parameters studied. Thus from these results dermatoglyphics can serve as an adjunct method in screening both Down's syndrome to aid early detection and bring about early intervention. The study recommends dermatoglyphic as a diagnostic tool in the early screening of Down's syndrome patients in Nigeria.

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References

- Adeyokunnu, A. (1982). The incidence of Down's syndrome in Nigeria. Journal of Medical Genetics, 19: 277-279.
- Aminu, A.T., Taura, M.G. and Adamiu, L.H. (2014). Palmar creases, a comparative study between Epilepsy patients and healthy subjects among Hausas of Northern Nigeria. *American Journal of Medicine and Medical Sciences*, 4(5): 175-179.
- Arrieta, M.I., Martinez, B., Criado, B., Simon, A., Salazar, L. and Lostao, C.M. (1990). Dermatoglyphic analysis of Autistic Basque children. *American Journal of Medical Genetics*, 35(1): 1-9.
- Babler, W.J. (1991). Embrological development of epidermal ridges and their configuration. *Birth Defects Original Articles Series*, 27(2): 95-112.
- Barbosa, E.D., Fernardes, P.R. and Fernardes, F.J. (2009). Anthropology, muscular strength and dermtoglyphics of individuals with Down's syndrome. *Fitness and Performance Journal*, 8(4): 269-78.
- Boroffice, R.A. (1978). Down's's syndrome in Nigeria: Dermatoglyphic analysis of 50 cases. *Nigeria Medical Journal*, 8(6): 571-6.
- Bryant, J.I., Emmanuel, I., Huang, S. and Kronmal, R. (1970). Dermatoglyphs of Chinese children with Down's syndrome. *Journal of Medical Genetics*, 7: 338-344.
- Bulagouda, R.S., Patil, P.J., Hadimani, G.A., Bannur, B.M., Patil, B.G., Mallashetty, N.S. and Bagoji, I.B. (2013). Study of palmar dermatoglyphics in patients with essential hypertension between the age group of 20-50 years. *International Journal of Medical Research and Health Sciences*, 2(4): 773-777.
- Butler, M.G. and Meaney, F.J. (2005). *Genetics of Developmental Disabilities*, 123-124, Taylor and Francis Group, United State.
- Campbell, E.D. (1998). Fingerprints and palmer dermatoglyphics. *E-fingerprints.net*.
- Cummins, H. and Midlo, C. (1943). Finger prints, Palm and soles: An introduction to dermatoglyphic. 315, Blakiston, Philadelphia.
- Daniel, W.W. (1999). *Biostatistics: A Foundation for Analysis in the Health Sciences* (7th Ed). John Wiley & Sons, New York.
- Fogle, T. (1990). Using dermatoglyphics from Down's syndrome and class populations to study the genetics of a complex trait. in Tested studies for laboratory teaching (Ed. C.A. Goldman), *Proceedings of the Eleventh workshop/conference of the association for biology laboratory education*, 11: 129-150.
- Gordon, G., Peter, G., Paul, R. and Malcolm, R. (2010). Learning disability: A life cycle approach to valuing people. 43-44, McGraw-Hill International.
- Holt, S. (1968). Genetics of dermal ridges: The relation between total ridge count and variability of counts from finger to finger. *Galton Laboratory*, 22: 323-339.
- Jeewandeep, K. and Arvinder, P.S.B. (2013). Role of dermatoglyphics in medical disorders. *Indian Journal of Fundamental and Applied Life Science*, 3(3): 536-539.
- Johnny, F. (2018). The development of the study of dermatoglyphics. Retrieved 20/01/2018.
- Kiran, K., Kavitha, R. and Amitha, M.H. (2010). Dermatoglyphics as a non-invasive diagnostic tool in predicting mental retardation. *Journal of International Oral Health*, 2(1): 95-100.
- Lainhart, J.E., Piven, J., Wzorek, M., Landa, R., Santangelo, S.L., Coon, H. and Folstein, S.E. (1997). Macrocephaly in children and adults with autism. *Journal of American Academy and Child Adolescent Psychiatry*, 36: 282-290.
- Malla, T.M. and Srivastava, P.G.N. (2008). Dermatoglyphic variations among clinically diagnosed Down's syndrome cases: A cohort study. *Biomedical and Pharmacology Journal*, 1(2): 527-532.
- Milicic, J., Bujas, P.Z. and Bozikov, J. (2003). Dermatoglyphs of digito-palmar complex in autistic disorder: family analysis. *Croatia Medical Journal*, 44(4): 469-76.

- Mollic, M.J.H. and Habib, M.A. (2011). Dermatoglypichs a good tool in preventive medicine. *Journal of Armed Forces Medical College*, 7(2): 01-02.
- Moore, K.L. and Persaud, T.V.N. (2003). *The Developing Human: Clinical Oriented Embryology* (7th Edition), 486, Saunders, India.
- Oghenemavwe, L.E. and Osaat, R.S. (2015). An improvise easy digital method for palmar and plantar dermatoglyphics. *Bioscience and Bioengineering*, 1(3): 88-89.
- Oladipo, G.S. and Ogunnowo, B.M. (2004). Dermatoglyphic patterns in diabetes mellitus in a South Eastern Nigeria population. *African Journal of Applied Zoology and Environmental Biological*, 6: 6-8.
- Oladipo, G.S., Afolabi, E.O. and Esomonu, C. (2010). Dermatoglyphic patterns of obese versus normal weight Nigeria individuals. *Biomedicine International*, 1: 66-69.
- Oladipo, G.S., Dike, E.U. and Okoh, P.D. (2012). A comparative study of the digital pattern, position of Triradii, b-c and a-d palmar distances of diabetic subjects and essential hypertensive individuals in Rivers State. *International Journal of Advanced Biotechnology and Research*, 3(2): 615-620.
- Oladipo, G.S., Okoh, P.D., Oghenemavwe, L.E. and Yorkum, L.K. (2013). Dermatoglyphic patterns of Autistic children in Nigeria. *Journal of Biology, Agriculture and Healthcare*, 3(7): 80-83.
- Oladipo, G.S., Olotu, E.J., Fawehinmi, H.B., Okoh, P.D. and Iboroma, A.D. (2007). Dermatoglyphics in idiopathic (primary) dilated cardiomyopathy in South Southern Nigeria. *Scientific Research and Essay*, 2(10): 416-420.
- Oladipo, G.S., Paul, C.W., Bob-Manuel, I.F., Fawehinmi, H.B. and Edibamode, E.I. (2009). Study of digital and palmar dermatoglyphic patterns of Nigerian women with malignant mammary neoplasm. *Journal of Applied Biosciences*, 15: 829-834.
- Osaat, R.S., Didia, B.C., Osunwoke, E.A., Oladipo, G.S., and Victor, P.D. (2019). Dermal AB ridge count: Possible marker for Autism. *European Journal of Biomedical and Pharmaceutical Sciences*, 6(8): 92-96.
- Osaat, R.S., Oghenemavwe, E.L., Oladipo, G.S. and Amadi, P.N. (2022). Dermatoglyphics: A diagnostic tool for Autism. *Journal of Anatomical Sciences*, 13(1): 149-154.
- Ozyurt, B., Songur, A., Sarislmaz, M., Akyol, O., Namli, M. and Demorel, R. (2010). Dermatoglyphics as markers of prenatal disturbances in Schizophrenia: A case-control study. *Turkish Journal of Medical Sciences*, 40(6): 917-924.
- Plato, C.C., Cereghino, J.J. and Steinberg, F.S. (1973). Plamar dermatoglyphics of Down's syndrome: Revisted. *Pediatrics Research*, 7: 111-118.
- Pratibha, R., Abhilash, P.R., Herald, J.S., Anuja, N., Priya, P. Chandrasekar, T., Sentamilselvi, G. and Janaki, V.R. (2011). Conventional dermatoglyphics –revived concept: A review. *International Journal of Pharmacology and Biological Sciences*, 2(3): 446-457.
- Pushpa, B., Kazi, S.N. and Vatsalaswamy, V.A. (2013). Role of dermatoglyphic fingertip patterns in the predition of maturity onset diabetes mellitus (Type II). *Journal of Dental and Medical Sciences*, 8(1): 01-05.
- Raizada, A., Johri, V., Ramnath, T., Chowdhemy, D.S. and Garg, R.P. (2013). A cross sectional study on palmardermatoglyphics in relation to carcinoma breast patients. *Journal of Clinical and Diagnosis Research*, 7(4): 609-612.
- Rajangam, S., Janakiram, S. and Thomas, I.M. (1995). Dermatoglyphics in Down's syndrome. *Journal of Indian Medical Association*, 93(1): 10-13.
- Ramesh, M., Kumari, K.G., Kalpana, V.L. and Sudhakar, G. (2012). Palmar and digital dermatoglyphic patterns in sickle cell anemia patients of North Coastal Andhra Pradesh, South India. *Antrocom Online Journal of Anthropology*, 8(1): 23-32.
- Reed, T., Sprague, F.R., Kang, K.W. and Nance, W.E. (1975). Genetic analysis of dermatoglyphic patterns in twins. *Human Heredity*, 25: 263-275.

- Reed, T., Uchida, A.I., Norton, J.A. and Christian, J.C. (1978). Comparisons of dermatoglyphic patterns in monochorionic and dichorionic monozygotic twins. *American Journal of Human Genetics*, 20: 383-391.
- Sandeep, V.P., Bharat, S.B., Megha, A.D. and Vigary, P.M. (2012). Study of the fingertip pattern as a tool for the identification of the dermatoglyphic tract in bronchial asthma. *Journal of Clinical and Diagnostic Research*, 6(8): 1397-1400.
- Schaumann, B. and Alter, M. (1976). Dermatoglyphics in medical disorders. *Newyork Springer Verlag*, 27-87, Berlin.
- Sharma, A., Sood, V., Singh, P. and Sharma, A. (2018). Dermatoglyphics: A review in fingerprints and their changing trends of use. *CHRISMED Journal of Health & Research*, 5(3): 167-172.
- Sharma, M.K., Jhawar, P., Sharma, H., Sharma, S. and Kalavatia, I. (2012). Dermatoglyphics an attempt to predict Down's syndrome. *International Journal of Biological & Medical Research*, 3(2): 1631-1635.
- Shield, J.P., Wadsworth, F.J.H. and Baum, J.D. (1995). Dermatoglyphics fetal growth and diabetes in children. *Archivesof Disease in Childhood*, 72: 159-160.
- Singh, A., Gupta, R., Zaidi, S. and Singh, A. (2016a). Dermatoglyphics: A brief review. *International Journal of Advanced and Integrated Medical Sciences*, 1(3): 111-115.
- Singh, S., Khurana, A.K., Harode, H.A., Tripathi, A., Pakhane, A. and Chaware, P. (2016b). Study of fingerprint patterns to evaluate the role of dermatglyphics in early detection of bronchial asthma. *Journal of Natural Science, Biology and Medicine*, 7(1): 43-46.
- Slatis, H.M., Katznelson, M.B. and Bonne-Tamir, B. (1976). The inheritance of fingerprint patterns. *American Journal of Human Genetics*, 28(3): 280-289.
- Stevenson, R.E., Hane, B., Arena, J.F., May, M., Lawrence, L., Lubs, H.A. and Schwartz, C.E. (1997). Arch finger prints, hypotonia and areflexia associated with x-linked mental retardation. *Journal of Medical Genetics*, 34(6): 465-469.
- Stosljevic, M. and Adamovic, M. (2013). Dermatoglyphic characteristics of digito-palmar complex in autistic boys in Serbia. *Vojnosanitetski Pregled*, 70(4): 386-390.
- Tarca, A. (2008). New dermatoglyphic investigations on infantile autism. *Journal of Preventive Medicine*, 16(1-2): 69-76.
- Tarca, A. and Barabolski, C. (2003). Pathology of deermatoglyphics in infantile autism. *The Journal of Preventive Medicine*, 11(1): 11-17.
- The Secret of Your Child Fingerprints (2011). http://dermatoglyphics.org
- Verbov, J. (1970). Clinical significance and genetics of epidermal ridges A review of dermatoglyphics. *Journal of Investigative Dermatology*, 54, 261-271.
- Villar, A.J. and Epstein, C.J. (2005). Down's syndrome. *Encyclopedia of Life Sciences*, John Wiley & Sons Ltd., www.els.net
- Walker, H.A. (1977). A dermatoglyphic study of autistic patients. *Journal of Autism and Childhood Schizophrenia*, 7(1): 11-21.