# Dermatoglyphic Patterns in Cystic Fibrosis Children

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#### Abstract

*Objective:* It is believed that fingerprints and palm patterns may represent genetically determined congenital abnormalities in Cystic Fibrosis (CF). The main idea of this paper was to determine differences of fingerprints and palm patterns in CF and normal children.

*Methods:* Forty-six CF children (27 males, 19 females) and 341 (113 males, 228 females) healthy individuals were recruited for this study. Fingerprint patterns, Total ridge count (TRC) of each finger, a-b ridge count, and atd angles of all participants were recorded. Asymmetry of the right and left hand for each value was determined and dissimilarity in fingerprint patterns between homologous fingers was compared using Chi-square analysis, Mann-Whitney U test and Fisher's exact test.

**Findings:** There were significant differences in the mean TRC of the right digit IV (P=0.009), left digit III (P=0.02), left digit IV (P=0.03), and left digit V (P=0.03). Furthermore, we found significant differences in right atd angel (P=0.001), left atd angel (P=0.002), right a-b ridge (P=0.007) and left a-b ridge (P=0.001). In contrast, we found no significant differences in atd angle asymmetry, a-b ridge count asymmetry and pattern dissimilarity score between both groups (P>0.05).

*Conclusion:* Dermatoglyphic characteristics could be used as a supplementary diagnostic method in CF children.

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Key Words: Dermatglyphics; Cystic Fibrosis; Palm Patterns; Fingerprints; Children

## **Introduction**

Dermatoglyphics is the science of fingerprints and palm prints analysis which are constant throughout life<sup>[1-3]</sup>. It is believed that they may represent different genetically determined congenital abnormalities. Several genes are involved in the inheritance of the dermal traits<sup>[1]</sup>. Dermal ridges form in 6<sup>th</sup> week of gestation and reach their maximum size between 12<sup>th</sup> and 13<sup>th</sup> week. Historically, dermatoglyphics was a classical model to consider the polygenic inheritance. Therefore, some specific dermatoglyphic patterns may be accompanied with different genetic abnormalities<sup>[1,3]</sup>.

Recent studies on dermatoglyphics have shown especial patterns in various congenital abnormalities such as skeletal maturation, diabetes type 1, schizophrenia, albinism, cleft lip and palate, rheumatoid arthritis, congenital spinal cord anomalies and Klinefelter syndrome<sup>[4-10]</sup>.

Cystic Fibrosis (CF) is a common fatal,

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autosomal recessive disease in Caucasians<sup>[11,12]</sup>. CF is a progressive disease that involves exocrine glands, lungs, gastrointestinal system, pancreas, liver, kidneys and reproductive system<sup>[3,12,13]</sup>. CF prevalence varies from 1 to 2500 among Caucasians<sup>[13]</sup>; 10 million carriers of the recessive gene exist in the world<sup>[14]</sup>. CF can be diagnosed antenatally using genetic tests and in early childhood with sweat test<sup>[15]</sup>.

The main idea of this paper was to determine the differences in dermatoglyphic patterns in CF and normal children.

#### Subjects and Methods

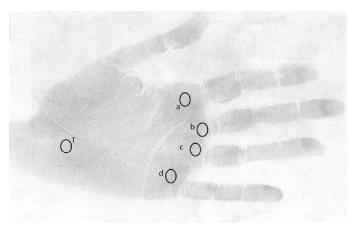
This is a case-control study performed between October 2012 and March 2013. The study group included 46 children with CF, confirmed by two positive sweat tests, referred to CF clinic in Dr. Sheikh Pediatric Hospital, Mashhad University of Medical Sciences, Iran. The control group consisted of 341 healthy participants. Exclusion criteria included children less than one year of age, individuals with current atopic dermatitis and any finger or palm abnormalities which interfered with the procedure. All subjects were informed about the process and consent forms were obtained. We used red powder blush to roll finger and palm separately. The powder was rubbed on palm and fingers of both hands by means of a brush. A piece of adhesive tape was placed on a rolling plate and the participant pressed his/her palm on the tape in distal to proximal direction. It was then transformed onto a labeled paper indicating right and left side. The same process was carried out for fingerprints as each finger was pressed separately to the adhesive tape. Fingers were named with roman numbers from I (thumb) to V (little finger).

Fingerprint pattern types were categorized into three groups, arch (A), loop (L) and whorl (W). Whorl was divided into two subgroups named simple whorl (W) and double whorl (2W) (Fig. 1). Patterns of homolog fingers in right and left hand were given 0 or 1 score due to their similarity or dissimilarity. Finally, these scores were added and final scores ranged 0 to 5<sup>[2]</sup>.

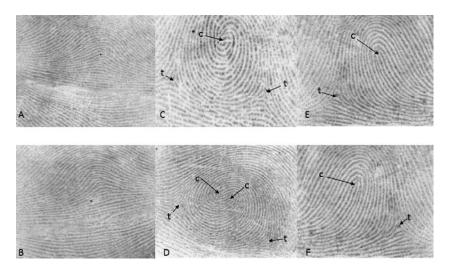
After the center point of each fingerprint was joined to triradius (i.e. the conjunction point of three opposing ridge systems), TRC was calculated by summing the total numbers of ridges between the two centers<sup>[16]</sup>.

Total ridge count of arch pattern is always zero because it has no triradius. Loop pattern has one triradius, but whorl pattern has two triradii. The higher calculated number of lines was reported for whorl pattern TRC (Fig. 1).

On the following step, we found the triradii (a, b, c, d) at the base of each finger (Fig. 2). a-b line



**Fig. 1:** Palm patterns are labeled a, b, c, d and t a: triradius "a" at the base of the 2<sup>nd</sup> finger b: triradius "b" at the base of the 3<sup>rd</sup> finger c: triradius "c" at the base of the 4<sup>th</sup> finger d: triradius "d" at the base of the 5<sup>th</sup> finger t: triradius "t" at the proximal part of the palm



**Fig. 2:** Fingerprint pattern types are categorized into three groups: arch (A,B), loop (F,G) and whorl. Whorls are divided into two subgroups named simple whorl (C) and double whorl (D). c: center point of fingerprint, t: triradius

was formed by joining a to b triradius. The ridges on the a-b line in both hands were calculated with magnifying glass.

After detection of triradius a (at the base of  $2^{nd}$  finger on palm print), t (on the proximal of palm) and d (at the base of  $5^{th}$  finger on the palm), atd angle was drawn and measured by protractor in both hands.

All the data was collected twice by one person and the mean of these two numbers was finally reported.

We used chi-square test for comparing frequency of pattern types, Fisher's exact test for comparing the fingerprint pattern types and the disease phenotype and Mann-Whitney test to compare a-b ridge count, TRC, atd angle and asymmetry of right and left hand between control and study groups. *P*-values less than 0.05 were considered statistically significant.

### **Findings**

A total of 387 participants, 46 CF and 341 healthy children were enrolled in this study. Table 1 shows the baseline characteristics of studied group.

Number and percentage of disease phenotype(failure to thrive, steatorrhea, clubbing, liver disease, other gastrointestinal symptoms and respiratory symptoms) and dermatoglyphic pattern types of the studied group are shown in Table 2. There was no significant difference in dermatoglyphic pattern types and phenotype.

Variable		Mean (SD)	Variable	n (%)
Age (month)		105.9 (43.0)	Relative parents	30 (85.7)
Weight		22.49 (8.8)	FTT	16 (57.1)
Height (cm)		116.4 (20.0)	Steatorrhea	13 (46.4)
HC <sup>a</sup> (cm)		50.98 (1.9)	Clubbing	1 (3.6)
MAC <sup>b</sup> (cm)		16.4 (2.3)	Respiratory symptoms <sup>c</sup>	15 (53.6)
Sweat test	Cl Na	101.7 (38.8) 104.7 (47.6)	Other gastrointestinal symptoms <sup>d</sup>	17 (60.7)
Age of onset (	month)	3.42 (6.29)	Liver disease	2 (7.1)
Age at diagnos	sis (month)	11.8 (17.2)		

## Table 1: Baseline characteristics of the studied groups

a:Head Circumference

b: Mid arm circumference

c: Other gastrointestinal symptoms include vomiting, GER, rectal prolapse, chronic diarrhea

d: Respiratory symptoms include pneumonia, cough, wheezing, dyspnea

611

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Digit	Pattern	FTT		Steatorrhea	ea	Clubbing		Liver disease	ase	Other		Respiratory	ĥ.
		Right hand n (%)	Left hand n (%)	Right hand n (%)	ght hand Left hand (%) n (%)	Right hand Left hand n (%) n (%)	Left hand n (%)	Right hand Left hand n (%) n (%)	Left hand n (%)	Right handLeft hand n (%) n( %)		Right hand n (%)	Left hand n (%)
	Loop	6 (60)	9 (75)	8(53.3)	6 (50)	(0)0	1(100)	1(6.7)	(0)0	10(66.7)	5(41.7)	5(33.3)	5 (41.7)
-	Arch		0 (0)		0 (0)	I	0(0)	I	0()0	I	(0)0	I	1 (100)
-	Whorl	7 (53.8)	7 (46.7)	5 (38.5)	7 (46.7)	1(100)	0(0)	1(7.7)	2(13.3)	7(53.8)	12(80)	10(76.9)	6 (60)
	<i>P.</i> value	0.7	0.17	0.43	1	0.46	0.46	1	0.52	0.7	0.05	0.05	0.4
	Loop	5 (50)	8 (57.1)	4 (40)	7 (50)	(0)0	1(7.1)	1(10)	1(7.1)	6(60)	9(64.3)	6(60)	6 (42.9)
I	Arch	2 (100)	2 (66.7)	1 (50)	1(33.3)	0(0)	0(0)	0(0)	0(0)	0(0)	0(0)	1(50)	2 (66.7)
1	Whorl	9 (56.3)	6 (54.5)	8 (50)	5(45.5)	1(6.3)	0(0)	1(6.3)	1(9.1)	11(68.8)	8(72.7)	8(50)	7 (63.6)
	<i>P.</i> value	0.5	1	0.84	1	1	1	1	1		0.05	0.8	0.5
	Loop	12 (63.2)	9 (52.9)	8 (42.1)	6(35.3)	1(100)	1(100)	2(10.5)	2(11.8)	12(63.2)	10(58.8)	9(47.4)	7 (41.2)
I	Arch	2 (100)	3 (75.0)	1 (50)	2(50)	0(0)	0(0)	0(0)	0()0	0(0)	1(25)	1(50)	3 (75)
I	Whorl	2 (28.6)	4 (57.1)	4 (57.1)	5(71.4)	0(0)	0(0)	0(0)	0()0	5(71.4)	6(85.7)	5(71.4)	5 (71.4)
	P. value	0.13	0.8	0.82	0.31	1	1	0.64	0.68	0.2	0.16	9.0	0.2
	Loop	8 (61.5)	7 (58.3)	6 (46.2)	4(33.3)	1(7.7)	1(8.3)	1(7.7)	1(8.3)	7(53.8)	7(58.3)	8(61.5)	7 (58.3)
M	Arch	1 (100)	1 (100)	0 (0)	0 (0)	0(0)	0(0)	0(0)	0(0)	0(0)	0()0	0(0)	0 (0)
2	Whorl	7 (50)	8 (53.3)	7 (50)	6 (60)	0(0)	0(0)	1(7.1)	1(6.7)	10(71.4)	10(66.7)	7(50)	8 (53.3)
	<i>P.</i> value	0.8	1	1	0.25	0.5	0.4	1	1	0.2	0.42	0.5	0.8
	Loop	9 (50)	8 (50	6 (33.3)	6(37.5)	1(100)	1(6.3)	2(11.1)	1(6.3)	11(61.1)	9(56.3)	10(55.6)	11 (68.8)
Λ	Arch		1 (100)		0()0	I	0(0)	I	(0)0	I	(0)0	ł	0 (0)
	Whorl	7 (70)	7 (70)	7 (70)	7(70)	0(0)	0(0)	0(0)	1(10)	6(60)	7(70)	5(10)	4 (40)
	<i>P.</i> value	0.4	0.53	0.11	0.16	1	1	0.52	1	1	0.4	1	0.1
a: other { b: respira	gastrointestina atory sympton	a: other gastrointestinal symptoms include vomiting, GER, rectal, Chronic b: respiratory symptoms include Pneumonia, Cough, wheezing, dyspnea	e vomiting, GI nia, Cough, w		ectal, Chronic diarrhea zing, dyspnea	a							

Table 2. Comparing the number and percentage of disease phenotype and dermatoglyphic pattern types in CF patients

Digit	pattern type	CF group n (%)	Right hand Control n (%)	Total (%)	CF group n (%)	Left hand Control n (%)	Total (%)
I	Loop Arch Whorl P. value	25 (54.3) 2 (4.3) 19 (41.3)	162 (48.1) 8 (2.4) 167 (49.6) 0.5	187 (48.8) 10 (2.6) 186 (48.6)	24 (52.2) 2 (4.3) 20 (43.5)	173 (52.0) 17 (5.1) 143 (42.9) 1	197 (52.0) 19 (5.0) 163 (43.0)
II	Loop Arch Whorl <i>P.</i> value	21 (45.7) 5 (10.9) 20 (43.5)	162 (48.1) 29 (8.6) 146 (43.3) 0.9	183 (47.8) 34 (8.9) 166 (43.3)	24 (52.2) 6 (13.0) 16 (34.8)	159 (47.7) 28 (8.4) 146 (43.8) 0.4	183 (48.3) 34 (9.0) 162 (42.7)
III	Loop Arch Whorl <i>P</i> . value	32 (69.6) 4 (8.7) 10 (21.7)	222 (65.9) 24 (7.1) 91 (27.0) 0.7	254 (66.3) 28 (7.3) 101 (26.4)	29 (63.0) 6 (13.0) 11 (23.9)	212 (63.5) 27 (8.1) 95 (28.4) 0.5	241 (63.4) 33 (8.7) 106 (27.9)
IV	Loop Arch Whorl <i>P</i> . value	22 (47.8) 1 (2.2) 23 (50.0)	139 (41.6) 9 (2.7) 186 (55.7) 0.8	161 (42.4) 10 (2.6) 209 (55.0)	20 (43.5) 3 (6.5) 23 (50.0)	149 (45.2) 15 (4.5) 166 (50.3) 0.9	169 (44.9) 18 (4.8) 189 (50.3)
v	Loop Arch Whorl <i>P.</i> value	30 (65.2) 0 (0) 16 (34.8)	246 (74.3) 5 (1.5) 80 (24.2) 0.1	276 (73.2) 5 (1.3) 96 (25.5)	29 (64.4) 1 (2.2) 15 (33.3)	251 (77.2) 6 (1.8) 68 (20.9) 0.2	280 (75.7) 7 (1.9) 83 (22.4)

Table 3: Number and percentage of dermatoglyphic patterns in CF patients and control group

Table 3 shows number and percentage of fingerprint pattern types in case and control groups. There was no significant difference in fingerprint patterns of both groups.

Table 4 represents mean TRC in case and control groups. Significant differences in mean TRC between case and control groups were found in right digit IV (P=0.009), left digit III (P=0.02), left digit IV (P=0.03) and left digit V (P=0.03).

The TRC asymmetry for all digits is shown in Table 5. There was no significant difference

between both groups regarding TRC asymmetry.

According to Table 6, significant differences were found in right hand atd angle (P=0.001), left hand atd angle (P=0.002), right hand a-b ridge (P=0.007) and left hand a-b ridge (P=0.001) between case and control group.

No significant differences were found in atd angle asymmetry, a-b ridge count asymmetry or pattern dissimilarity score between both groups (Table 7). *P*-values less than 0.05 were considered statistically significant.

Digit		CF patients Mean Rank	Control group Mean Rank	<i>P</i> . value
т	R	156.85	197.35	0.2
1	L	181.49	191.74	0.5
II	R	156.43	196.30	0.2
11	L	181.49	191.74	0.5
III	R	167.84	194.15	0.1
111	L	153.87	195.54	0.02
11/	R	150.29	195.48	0.009
IV	L	155.51	192.54	0.03
v	R	168.20	191.88	0.2
v	L	152.44	188.30	0.03

## **Table 4:** Mean TRC of CF patients and control group

Digit	CF Patients Mean (SD)	Control group Mean (SD)	P. value
Ι	193.49	190.09	0.8
II	206.29	187.17	0.3
III	178.92	190.40	0.5
IV	175.41	188.63	0.4
V	174.02	183.10	0.6

Table 5: Total ridge count asymmetry for five digits in CF patients and control group

CF: Cystic Fibrosis; SD: Standard deviation

## **Discussion**

In this study, we observed significant differences in dermatoglyphic patterns including the mean TRC of the right digit IV, left digit III, left digit IV, left digit V and atd angle and a-b ridge in right and left hands of children with CF compared to control group.

Kobylisky et al reported significant differences in fingerprint pattern types <sup>[17]</sup>. They showed that arches fingerprint patterns were higher in CF females in contrast to higher loop patterns in CF males. Our results did not establish significant differences considering these patterns in both groups. Based on Weizman et al, whorl patterns were more frequent than loop patterns as opposed to control group in celiac patients<sup>[18,19]</sup>. In another study, whorl pattern values were also higher in celiac children and there was a correlation between dermatoglyphic patterns of celiac

Table 6: Mean (SD) for atd angles, a-b ridge counts in CF cases and control group

Group	CF cases	Control group	<i>P</i> -value
atd – R	234.19	179.94	$0.001^{**}$
atd - L	231.91	178.48	$0.002^{*}$
a-b-R	148.75	194.59	$0.007^{*}$
a-b-L	137.59	194.87	$0.001^{*}$

SD: Standard deviation; CF: Cystic Fibrosis

patients and their parents<sup>[21]</sup>. In 1986, Gottlieb et al observed that the arch pattern values were significantly higher in congenital syndrome of early onset constipation and abdominal pain<sup>[20]</sup>. Mathew et al found increased loop patterns in children with oral cleft<sup>[1]</sup>. In this study, we observed significant differences in the mean TRC of the right digit IV, left digit III, left digit IV, left digit V of CF cases compared to the control group.

compared to control group<sup>[3]</sup>. This finding was in contrast to Kobylisky et al

study<sup>[17]</sup> which reported no significant difference in the mean TRC. Kobylisky et al also showed that mean TRC values were lower in the CF group. Their results are in line with ours<sup>[17]</sup>. In 2002, Neiswanger et al observed no significant differences in TRC asymmetry in patients with

the mean TRC of the right digit IV, right digit V and

left digit II in patients with cleft lip (CLP)

Eslami et al observed significant differences in

 Table 7: Mean atd angle asymmetry, a-b ridge count asymmetry and pattern dissimilarity score in Cystic Fibrosis cases and control group

Group	CF cases	Control group	<i>P</i> -value
atd asymmetry	196.80	180.53	0.333
a-b asymmetry	183.54	186.34	0.8700
Pattern dissimilarity score	192.93	182.75	0.529

CF: Cystic Fibrosis

cleft lip who had a negative family history<sup>[21]</sup>. Their results were similar to those of our study.

Comparing the a-b ridge count, we found that a-b ridge count values were lower in CF patients. It was similar to Kobylisky et al study<sup>[17]</sup>. Rezaeinezhad et al showed that mean a-b ridge count values in patients with type 1 diabetes mellitus were higher in control group<sup>[10]</sup>.

In 1973, Taussing et al observed increased atd angle in CF children, which was in agreement with our results<sup>[22]</sup>. On the contrary, Kobylisky et al reported that the value of atd angles were significantly lower in CF females and significantly higher in CF males<sup>[17]</sup>. According to the study of Mathew et al, atd angles showed an increase in children with oral cleft<sup>[1]</sup>.

Esalmi et al observed that atd angles were not significantly different in CLP patients and control group. In 2013, Eslami et al observed no significant differences in a-b ridge asymmetry and pattern dissimilarity score in CLP patients, which was in agreement with our results <sup>[3]</sup>.

This study had its limitations. The limitations were confined to the small number of patients that may impose a negative effect on the final results. The authors recommend that further researches should be done in parents of CF children to assess the child and parent dermatoglyphic traits relation. Moreover, evaluating the distribution of fingerprint minutiae and palmar sweat glands in CF children would provide more comprehensive information of dermatoglyphic patterns in CF children.

## **Conclusion**

Dermatoglyphic characteristics were significantly different in CF children and control group. These traits could be used as a supplementary diagnostic method in CF children.

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#### Authors' Contribution

A. Ezzati: Concept, Acquisition of Data, Data Analysis, Manuscript Preparation, Critical Revision of the Manuscript

F. Batoeh: Prepared the design of the study and drafted and Approval of the Article

S.A. Jafari: Performed the study, acquired the data M.A. Kiani : performed the study, acquired the data N. Mahdavi-Shahri: Performed the study, acquired the data

H. Ahanchian: Performed the study, acquired the data S. Tehraniam: Manuscript Preparation, Critical Revision of the Manuscript

A. Jahanbin: Concept, acquired the data and Critical Revision of the Manuscript

H.R. Kianifar: Concept, prepared design of the manuscript, data analysis, and interpretation All authors approved final version of the paper.

#### Conflict of Interest: None

#### **References**

- 1. Mathew L, Hegde AM, Rai K. Dermatoglyphic peculiarities in children with oral clefts. *J Indian Soc Pedod Prev Dent* 2005;23(4):179-82.
- 2. Woolf CM, Gianas AD. Congenital cleft lip and fluctuating dermatoglyphic asymmetry. *Am J Hum Genet* 1976; 28(4):400-3.
- 3. Eslami N, Jahanbin A, Ezzati A. Palm and finger print characteristics in nonfamilial cleft lip and palate patients and their parents. *J Craniofac Surg* 2013; 24(3):769-72.
- Cam FS., Gul D, Tusuf Y, et al. Analysis of the dermatoglyphics in Turkish patients with Klinefelter's syndrome. *Hereditas* 2008;143:163-6.
- 5. Rajangam S, Ravindranth R, Shubha R, et al. Dermatoglyphics-quantitative analysis in rheumatoid arthritis. *Anthropologist* 2008;10(3): 233-5.
- 6. Jahanbin A, Mahdavishahri N, Naseri MM, et al. Dermatoglyphic analysis in parents with nonfamilial bilateral cleft lip and palate children. *Cleft Palate Craniofac J* 2010; 47(1):9-14.
- Vidal-Linhares R, De Oliveira-Matta M, Perrout-Lima J, et al. The relationship between adolescents' dermatoglyphic characteristics and skeletal maturation. *Rev Salud Publica (Bogota)* 2010;12(6): 929-37.
- Shakibaei F, Asadollahi GA, Tabibi A. Dermatoglyphics in patients with schizophrenia. J Res Med Sci 2011; 16(8):1055-61.
- Ghodsi Z, Mahdavi Shahri N, Khajeh Ahmadi S. Quantitative and qualitative study of dermatoglyphic patterns in albinism. *Curr Res J Biological Sci* 2012;4(4):385-8.

- 10. Mahdavii Shahri N, Rezaeinezhad H. Application of dermatoglyphic traits for diagnosis of diabetic type 1 patients. *Int J Environm Sci Developm* 2010;1:36-9.
- Nussbaum R, McInnes RR, Willard HF. Thompson & Thompson Genetics in Medicine. 7<sup>th</sup> ed. Philadelphia: Elsevier Health Sciences. 2007.
- Kianifar H-R, Bakhshoodeh B, Hebrani P, et al. Qulaity of life in cystic fibrosis children. *Iran J Pediatr* 2013;23(2):149-53.
- 13. Farjadian S, Moghtaderi M, Kashef S, et al. Clinical and genetic features in patients with cystic fibrosis in Southwestern Iran. *Iran J Pediatr* 2013;23(2): 212-5.
- 14. Modi AC, Quittner AL. Validation of a diseasespecific measure of health-related quality of life for children with cystic fibrosis. *J Pediatr Psychol* 2003; 28(8):535-46.
- 15. Hakkak AM, Keramatipour M, Talebi S, et al. Analysis of CFTR gene mutations in children with cystic fibrosis, first report from North-East of Iran. *Iran J Basic Med Sci* 2013; 16(8):917-21.
- 16. Durham N, Plato C. Trends in Dermatoglyphic Research. 1st ed. Amsterdam: Kluwer Academic Publishers. 1990.

- 17. Kobyliansky E, Bejerani M, Yakovenko K. Relationship between genetic anomalies of different levels and deviations in traits. Part 5: Dermtoglyphic peculiarities of males and females with cyctic fibrosis. *Z Morphol Anthropol* 1999;81: 249-302.
- Weizman Z, Vardi O, Binsztok M. Dermatoglyphic (fingerprint) patterns in celiac disease. J Pediatr Gastroenterol Nutr 1990; 10(4):451-3.
- 19. Tahan S, Medeiros EH, Wehba J. Dermatoglyphic patterns in celiac disease. *Arq Gastroenterol* 1997; 34(3):196-204.
- 20. Gottlieb SH, Schuster MM. Dermatoglyphic (fingerprint) evidence for a congenital syndrome of early onset constipation and abdominal pain. *Gastroenterology* 1986; 91(2):428-32.
- 21. Neiswanger K, Cooper M, Weinberg S, et al. Cleft lip with or without cleft palate and dermatoglyphic asymmetry: evaluation of a Chinese population. *Orthodon Craniofac Res* 2002;5(3):140-6.
- 22. Taussig L, Braunstein G, White BJ, et al. Silver-Russell dwarfism and cystic fibrosis in a twin: endocrine, chromosomal, dermatoglyphic, and craniofacial studies. *Am J Dis Child* 1973; 125(4):495-503.