Review Article

DOI: 10.5455/2320-6012.ijrms20140207

Dermatoglyphics: in health and disease - a review

Gh. Mohd. Bhat*, M. Arif Mukhdoomi, Bahir Ahmed Shah, Mohd Saleem Ittoo

Department of Anatomy, Govt. Medical College, Srinagar - 190001, J & K, India

Received: 12 November 2013 Accepted: 2 December 2013

***Correspondence:** Dr. Gh. Mohd. Bhat, E-mail: gmbhat144@gmail.com

© 2014 Bhat GM et al. This is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

ABSTRACT

Dermatoglyphics is the study of finger print patterns and the term was coined by Harold Cummins in 1926. Finger prints are imprints of epidermal ridges, which are formed in early embryonic life, during 10th to 16th week of intrauterine life and remain permanent during whole life. Dermatoglyphic patterns have polygenic inheritance and are affected by environmental factors in the uterus. Finger print patterns are mainly of three types: arches whorls and loops; though there are more than 100 ridge characteristics, called Galton's details, in a single rolled finger print. Dermatoglyphics is not only used in the identification of an individual but also serves as a mirror of one's potential and talent. In this review, we will be discussing Dermatoglyphics and its important role in the diagnosis of chromosomal disorders and other diseases which have some genetic bases.

Keywords: Dermatoglyphics, Finger prints, Arch, Whorl, Loop

INTRODUCTION

Dermatoglyphics (from Greek: derma= skin, glyph= carving) is the scientific study of the skin ridge patterns on the fingers, toes, palms of hands and soles of feet. The purpose of these ridges is to impart firmer grip and to avoid slippage. Dermatoglyphics can be traced back to 1892, when one of the most original biologists of his time Sir Francis Galton, a cousin of Charles Darwin, published his work on fingerprints. The study was latter on termed as Dermatoglyphics by Dr. Harold Cummins, even though the process of finger print identification had already in use for several hundred years.¹

The most famous of ancient finger print designs are carvings on the wall of Neothalic burial passage, situated on the island of Brittany L'iie de Gavr'inis, its inner walls are covered with incised designs of circular patterns, spirals, arches, sinuous and straight lines occurring on various combinations.² In ancient time, finger prints were used on pottery to indicate the maker

and brand of pottery. In ancient Assyria, finger prints served as a seal to give authenticity to documents of importance. They appeared in clay tablets, now treasured in British Museum.³ An aboriginal Indian carving was found at the edge of Kejimkoojir lake in Novascotia, where within the outlines of human hand carved in stone showed lines which represented dermatoglyphics and flexions creases.¹ Sir William Herschel, a collector in Bengal (1858) was the first person to take finger and hand prints of a contractor which is still preserved in museum of London. He used the system for identification of criminals.³

The Ridges are differentiated in their definitive form during third to fourth month of fetal life and once formed remain permanent and never change throughout the life except in dimensions in proportion to the growth of an individual. The original ridge characteristics are not disturbed unless skin is damaged to the depth of 1mm.³ Development of dermatoglyphic pattern is under genetic control. This is evident from the clear resemblance of dermatoglypics among related persons. There are many diseases known to be caused by abnormal genes. Whenever there is any abnormality in the genetic makeup of parents, it is inherited to the children and is reflected in dermatoglyphic pattern.⁴

Dermatoglyphics as a diagnostic aid is now well established in number of diseases which have strong hereditary basis and is employed as a method for screening for abnormal anomalies.⁵ Apart from its use in predicting the diagnosis of genetic disease; dermatoglyphics is also used in forensic medicine in individual identification, physical anthropology, human genetics and medicine. Sir Galton Francis (1892) published a book "finger prints", which included the first classification of finger prints. He used ridge characteristics called 'minutea', still called Galton's details.⁶

DEVELOPMENT OF EPIDERMAL RIDGES

The ridge pattern depends upon the cornified layer of epithelium and dermal pattern. There is proliferation of cells in the lower zone of epidermis which projects in the dermis as regular spaced thickenings and the dermis subsequently projects upwards in the epidermal hollows, called dermal papillae. This is followed by appearance of elevations formed by them on the skin surface which are known as epidermal ridges.¹

Differentiation of epidermal ridges takes place early in fetal development. The ridge pattern is genetically determined and is affected by environmental factors. There exists relationship between epidermal ridge and fetal volar pads, because in course of development the ridge pattern is formed at the site of these pads.¹ Environmental factors such as external pressure on fetal pads and embryonic fetal finger movements could influence ridge formation.⁸ Dermatoglyphics traits such as such as finger ridge count develop between 10th to 17th week post conception.⁹ Dermatoglyphic features are inherited through a polygenic system with individual genes contributing an additive genetic component.8, ¹⁰Ridge formation is influenced by individual differences in developmental stability and first and second trimester insults on the embryo result in dermatoglyphic changes.8,11

FINGER PRINT TYPES

There are three types of finger prints:

- 1. Visible prints are also called as patent prints because these are visible to the naked eye without development and are left in some medium like blood, dirt, ink or grease on the finger.
- 2. Latent prints are not apparent to the naked eye and are formed from water, salt, amino acids and oils

contained in the sweat. These can be made visible by dusting, fuming or chemical reagents.

3 Impressed prints or plastic prints. These prints are indentations left on soft pliable surfaces such as clay, wax, paint or another surface that will take the impression. These are visible to the naked eye.

PATTERNS OF FINGER PRINTS

There are three basic finger print patterns a) Arches b) Loops c) Whorls¹²

- a) Arches: These are found in five percent of finger print patterns. The ridges run from one side to another of patterns, making no backward turns. Ordinarily there is no delta in an arch pattern. There are four types of arch patterns 1) Plain arches 2) Radial arches 3) Ulnar arches 4) Tented arches.¹³
- b) Loops: Loops occurs in about 60 to 70% of finger prints. One or more ridges enters on either side of impression, recurves, touches or crosses the line running from the delta to the core and terminates on or in the direction of the side ,where the ridge or ridges entered. In ulnar loop the ridges open on the ulnar side, in radial loops the ridges open on the radial side.¹³
- Whorls: These are seen in about 25 to 35 % of finger c) print patterns. In a whorl some of the ridges make a turn through at least one circuit. Any finger print pattern which contains two or more deltas will be a whorl pattern. There are six types of whorls: a) concentric whorl - the ridges are arranged in concentric rings around the core. b) Spiral whorl the ridges spiral around the core in clockwise or anticlockwise direction. c) Mixed whorl- it contains circles and spirals in same pattern. d) Central pocket whorl-it contains a smaller pocket within a loop. e) Twin whorl-in these ridges arising from each core open towards the opposite margin of the finger. f) Accidental whorls-these represents combination of two or more of above configurations.¹³

RIDGE CHARACTERISTICS

A single rolled finger print may have as many as hundred or more identification points, called as ridge characteristics which are as follows:^{8,14,15}

- 1. Ridge dots an isolated ridge unit whose length approximates its width in size.
- 2. Bifurcation the point at which one friction ridge divides into two friction ridges. It can be double bifurcation or opposite bifurcation.
- 3. Trifurcation the point at which one friction ridge divides into three friction ridges.

- 4. Ending ridge a single bifurcation ridge that terminates within the friction ridge structure.
- 5. Ridge crossing a point where two ridge units intersect.
- 6. Enclosure a single friction ridge that bifurcates and rejoins after a short course and continues as a single friction ridge.
- 7. Spur a bifurcation with one short ridge branching of a longer ridge.
- 8. Bridge a connecting friction ridge between parallel running ridges.

OTHER LAND MARKS

- a) Triradii a triradius is formed by the confluence of three ridge systems.
- b) Core it is approximate center of the palm. The core may be of different shapes. In ridge counting the point of core (not the whole core) is used.
- c) Radiants these are ridges that arise from triradius and enclose the pattern area.
- d) Delta the point on a ridge, at or in front of and nearest the centre of divergence of the type lines delta area is a triangular area from where the ridges radiate outwards.

Animal finger prints: Humans are not the only ones with the finger prints. Some primates like gorillas and chimpanzees and kola bear have their own unique finger prints.

Adermatoglyphia: It is a rare condition which causes people to be born without any finger prints. It is also known as immigration delay, disease because affected individuals report significant difficulties on entering countries that requires finger print recording.^{16,17}

DERMATOGLYPHICS IN HEALTH

The type of finger print is unique based on the genetic characteristics of each individual. In addition of predictive value of finger prints in various diseases dermatoglyphics is used in identification of an individual. Identification is a set of physical characteristics, functional or psychic, normal or pathological that defines an individual.¹⁸ Due to uniqueness of finger prints , these can be used to identify the criminals at crime scene, dead or unconscious person in blast injuries or mass disaster injuries , accidental exchange of new born babies, in prevention of impersonation of cheques, bank notes and even for national identification.¹⁹ Dermatoglyphics can be used to determine or exclude parentage under circumstances of uncertainty of paternity of child.²⁰

Dermatoglyphics is like a map that allows one to understand his own potential and talent. Temperament and character can be correlated with finger print pattern. Whorl signifies stubbornness, composite is a sign of faithless and unreliable character, loop signifies lack of perservance, arch denotes merciless crude behavior. A person with ulnar loop on all fingers is clear spirited, mild mannered and strong willed person (melancholic), cool in judgment and ruthless in business. A person having whorls on all fingers is restless, doubting, sensitive, clever, eager for action and inclined to crime. A mixture of whorls and loops signifies a neutral character, kind, obedient, truthful but often undecided and impatient. Arches and radial loops occurs in person who is ambitious, cool, stubborn, disobedient, defiant and rebellious.20

There is a correlation between cephalic size, form of an individual and type of finger print pattern. In Chinese (brachycephalic) there is increased frequency of whorls and arches, in English (dolichocephalic) there is reduction of whorls and increase in arches. There is association between whorls and blood group B, a loop and blood group A, person with blood group O have more loops and fewer whorls. In general females have narrow ridges, more arches and fewer whorls. Females also have large frequency of hypothenar IV interdigital patterns. The finger prints of imbecile and idiots are similar to the finger prints of monkeys. In these people palmar hypothenar pattern is dominant, arches are more, axial triradius located centrally and simian crease is present. In imbecile persons there is great reduction of whorls in the right index and ring fingers.²⁰

Brown Caucasoid and Indians showed higher frequency of patters in hypothenar area. Africans and Mongolians showed highest frequency in the fourth interdigital pattern, thenar and first interdigital pattern in Americans, second interdigital pattern in Nigroes and third interdigital pattern in Europeans. Dermatoglyphics has been used to differentiate between mono and dizygotic twins, which are not differentiable from DNA finger printing.²¹

DERMATOGLYPHICS IN DISEASE

Dermotoglyphics as a diagnostic tool is now well established in a number of diseases which have strong hereditary basis.^{22,23} The dermatoglyphic patterns are analyzed in various ways like a) Quantitative analysis of finger prints i.e. loops, arches, whorls. b) Total finger ridge count. c) Absolute finger ridge count. d) Position of axial triradii. e) Total number of palmer triradii. f) a-b ridge count g) ATD angle.^{8,24}

Commins H $(1936)^{25}$ was the first person to show the possible use of dermatoglyphics in clinical medicine. In the recent decades a considerable improvement has been achieved in the concept of relations between the types of fingers ridges and some individual disorders.²⁶⁻²⁸

Epidermal ridges are formed between 11-24 weeks of gestation and after this period epidermal ridges do not change.⁹ Since skin and brain develop from same ectoderm, dermatoglyphic variations are informative for disturbances.²⁹ early developmental brain Dermatoglyphics is considered as a window of a congenital anomalies and is sensitive indicator of intra anomalies.30 uterine The current status of dermatoglyphics is such that the diagnosis of some illnesses can now be done on the basis finger prints alone.²⁹ The ridge malformations may be congenital or acquired. The congenital malformations of dermatoglyphics are of four types: (a) Ridge aplasia implies absence of pattern (b) Ridge hypoplasia implies reduced height of ridges (c) Ridge dissociation means breaking of ridge- inherited as autosomal dominant trait (d) Ridge of the end indicates the vertical ridges which run off the end of the finger prints- also inherited as autosomal dominant triat.31

A. Dermaoglyphics in diseases which are purely genetic disorders

- 1. Down's syndrome (trisomy 21). These patients have mainly ulnar loops, significantly different atd angle, single transverse palmer crease (simian line), lower ridge count along digital midlines.^{32,33}
- 2. Turner's syndrome. Predominance of whorls, although the pattern of frequency depends on the particular chromosomal abnormality.³⁴
- 3. Klinefelter's syndrome. Excess of arches on digit 1, more frequent ulnar loops on digit 2, over all fewer whorls, lower ridge count for loops and whorls and total reduction of finger ridge count.³⁵
- 4. Patau syndrome. Excess of arches on the finger tips and single palmar crease in 60% people.³⁵
- 5. Edwards's syndrome (trisomy 18). 6-10 Arches on finger tips and single palmar crease in 30% people.³⁵
- 6. In inborn blindness there are abnormal triradii and excess of arches on finger prints.³⁶
- Noonan syndrome. Increased frequency of whorls on finger tips and the axial triradius 't' and increased incidence of single transverse palmer crease.³⁷

B. Dermatoglyphics in other diseases which have some genetic background

1. Neurological diseases:

Since brain and skin develop from same ectoderm, dermatoglyphic variations are informative for early developmental brain disturbances.⁸

- Alzheimer's disease: There is increased frequency of ulnar loops and decreased frequency of whorls and arches, radial loop on the 4th and 5th digits are more frequent like in Down's syndrome.³⁸
- b) Schizophrenia: There is significant increase in whorls and decrease in loops in male Schizophrenics and there is significant reduction in arches all in patients.⁴⁹
- c) Cerebral palsy: There is increased frequency of arches, radial loops and whorl pattern and decreased pattern of ulnar loop, decrease in TFRC and 'ab' ridge count.⁴⁰
- Neurofibromatosis: There is increased frequency of central pocket whorls on the little fingers of both hands and left little finger shows increase in TFRC and 'ab' ridge count.⁴¹
- e) Epilepsy: These patients show significant changes in 'ab' ridge count, TFRC, AFRC, atd, tad, td angles, lateral deviation, C-line, finger ridge count and distal deviation.⁴²
- 2. Heart diseases
- a) Congenital heart diseases: There is overall increased incidence in hypothenar pattern with increase in atd angle. There is increased frequency in Sydney line in Ventricular septal defect and Tetrology of Fallot, distal displacement of axial triradius is increased in PDA.^{43, 44}
- b) Rheumatic heart disease: There is decreased frequency of arches in males and increased frequency of whorls in females. There is increased frequency of patterns in 3rd interdigital area in males and decreased 'td' ridge count, increased multiple axial triradius in females.⁴⁵
- c) Coronary heart disease: The etiology of coronary heart disease is multifactorial, with genetic predisposition having an important role. In these patients there is significant decrease in loops and increase in whorls in males. The palmer pattern is significantly decreased in thenar area in females, 3rd interdigital area in both males and females, increase in '4' palmar triradii in males and '5' palmar triradii in females and significant decrease in '6' palmer triradii in both sexes.²⁴

3. Diabetes mellitus

In type 1 DM there is increased frequency in whorls, and decreased ulnar loop, increased frequency of Sydney line, and increased incidences of arches in females.⁴⁶ In Maturity onset diabetes mellitus, there is decrease in mean value of TFRC, AFRC, increase in arches and decrease in whorls.⁴⁷

4. Cancer cervix

There is decrease in frequency of ulnar loops and increase in arches with increase in 'atd' angle, decrease in TFRC and decrease in 3rd interdigital palmer pattern.⁴⁸

5. Leprosy

There is high frequency of palmer pattern in thenar and 1st interdigital area on left palm, with slight increase in frequency of distal axial triradii.⁴⁹

6. Essential hypertension

There is increase in TFRC, decreased frequency of axial triradius 't' in right palm of females and 't and t' in right palm of male, decreased atd angle and absence of axial of triradii in 10% cases.⁵⁰

7. Bronchial Asthma

There is higher frequency of whorls and lower frequency of arch.⁵¹

8. Rheumatoid Arthritis

There is increase in arches and decrease in loops and whorls in males, where as in females there is increase in whorls and decrease in loops on the 1^{st} finger of both hands, with increase in arches on 3^{rd} digit and whorls on 4^{th} digit of left hand.⁵²

9. Tuberculosis

There is predominance of whorls and decrease in loop pattern; mean TFRC and AFRC are higher with narrower atd angle.⁵

10. Carcinoma breast

The mammary buds begin to develop during 6th week of intrauterine life and at the same time finger ridges also begin to develop. In carcinoma breast patients there are 6 or more whorls in the total finger pattern, increase in whorls in right ring and little fingers.⁵⁴

11. Sickle cell anemia

There is increased frequency of whorls and decreased frequency of ulnar loops and in some cases there are Sidney creases.⁵⁵

CONCLUSION

Dermatoglyphics is one of the oldest and still the most useful techniques available to mankind. Finger ridge patterns are formed due to underlying interlocking pattern of dermal papillae which produce overlying corresponding epidermal ridges. This pattern is unique to an individual and is a classic model of polygenic inheritance. Dermatoglyphics, as a means of identification, has been used by man from ancient times, but use of dermatoglyphic features in the diagnosis of various diseases has received attention from 17th century. Dermatoglyphics plays an important role in the diagnosis of chromosomal disorders and other diseases which have genetic background. Since dermal ridges develop during 6th-13th weeks of gestation, genetic message carried in the genome - normal or abnormal - is deciphered during this period and is reflected in dermatoglyphics. Many structures of the body like the brain, mammary glands, lips, alveoli, plate etc. develop during the same period as the finger ridges, abnormal developmental insults on these structures in uterus is likely to be reflected in the dermatoglyphic patterns.

From the above discussion we conclude that dermatoglyphics is a simple, inexpensive and bed side diagnostic aid for conditions of chromosomal aberrations and various inheritable diseases.

Funding: No funding sources Conflict of interest: None declared Ethical approval: Not required

REFERENCES

- 1. Commins H and Midlo C. Palmar and planter epidermal configuration (Dermatoglyphics) in European, Americans. Am J Phys-Anthropol. 1926;9:471-502.
- 2. Holt SB. Dermatoglyphic pattern, eds. Genitical variations in human populations. Pregarnon: oxford; 1961: PP791.
- 3. Commins H and Midlo C. Finger prints of palms and soles. An Introduction to dermatoglyphics. INC, New York: Dover pub; 1943.
- 4. Walker JFA. Sex linked recessive finger print pattern. J Hered. 1964;32:279-80.
- 5. Commins H and Midlo C. Finger prints of palms and soles. An Introduction to dermatoglyphics. INC, New York: Dover pub; 1961.
- 6. Galton F. Finger prints. Facsimile Ed. New York and London: Mac Millon; 1892.
- 7. Bonnevie K. Studies on papillary pattern of human fingers. J Genet. 1924;15:1-11.
- Schaumann B and Alter M. Dermatoglyphics in medical disorders. New York: Springer Verlog; 1976: 187-189.
- Babler WJ. Embryological development of epidermal ridges and their configuration. In: C.C Plato, RM Garuto and BA Shaumann, Eds. Dermatoglyphics: Science in transition. 2nd ed. New York: Wiley liss; 1991; 27: PP95-PP112.
- Chakraborty R. The role of heredity and environment on dermatoglyphics traits. In: C.C Plato, RM Garuto and BA Shaumann Eds. Dermatoglyphics: Science in transition. 2nd ed. New York: Wiley Liss; 1991; 27: PP151-PP192.

- 11. Marko TA. Genetics and developmental stability: an integrative conjecture on etiology and Nerobiology of Schizophrenia. Psychological Medicine. 1992;22:295-305.
- United states Department of Justice (FBI). The science of finger prints classification and uses. US government posting offices. 1984. Available at: http://bookstore.gpo.gov/products/sku/027-001-00033-5.
- Jalali F, Hajian Tilaki KO. A comparative study of dermatoglyphic patterns in patients with myocardial infarction and control group. Acta Medica Iranica. 2002;40(3):187-91.
- 14. Martin NG, Evas V, Loesch DZ. A genetical analysis of co variation between finger ridge counts. Am Hum Biol. 1982;9:539-52.
- 15. Bank SD, Pa DIP, Mukerjee DP. Finger dermayoglyphic variation in Rengma Nagas of Nagaland, India. Coll. Antropol. 2009;33:31-5.
- 16. Burger B, Fuchs D, Sprecher E, Itin P. The immigration delay desiese: adermatoglyphia inherited absence of epidermal ridges. J Am Acad dematol. 2011;64(5):979-80.
- 17. Nousbeck J, Burger B, Fuchs-Telem D, Parlovsky M, Fenig S, Sarig O, Itin P, Spercher E. A mutation in a skin specific isoform of SMARCADI causes autosomal dominant a dermtoglphia. Am J Hum Genet. 2011;89(2):302-7.
- 18. Rastogi P, Pillai KR. A study of finger prints in relation of gender and blood groups. J indian Acad Forensic Medicine. 2010;32:11-3.
- Hassan Q, Mustafa G, Yousufani, Ishaaq M, Abaasi MH. Comparative study of dermatoglyphics among the students of Ziauddin University. Med forum. 2011 Dec.;22(12):16-25.
- 20. Cummins H, Midlo C. Finger, palm and sole prints. An introduction to dermatoglyphics, Second ed. New York: Dova; 1943.
- 21. Uchida IA, Patau K, Smith DW. Dermal Pattern of 18 and D1 trisomies. Am J of Human Genetics. 1996;14(4);345-52.
- 22. Shiono H. Dermatoglyphics in Medicine. Am J Forensic Medicine Med Pathol. 1986;7(2):120-6.
- 23. Katz nelson M, Goldman B. Fetal dermatoglyphics. Clin Genet. 1982;21(4):237-42.
- 24. Chimne HD, Ksheersagar DD. Dermatoglyphics in Angiographically proven coronary artery disease. J Anat Soc India. 2012;61(2):262-8.
- 25. Commins H. Dermatoglyphics stigmata in Mangolisim. Anat Record. 1936;64(suppl.2):11.
- 26. Shamsuddin S, Masomi M, Magad Hossini M. Relations between the lines on the fingers of hands and the incidence of disease in the human. Journal of Kermin Medical Science University. 1997;4(3):136-42.
- 27. Simsek S, Taskiran H, Karakaya N. Dermatoglyphic analysis in children with CP. Neurobiology BP. 1998;6(3):373-80.

- 28. Verma SL, Chary TV, Singh S, Ashorom Z. Dermatoglyphic patterns in Schizophrenic patients. Acta Psychiatry-11. 1995;91(3):213-5.
- 29. Van O el CJ, Baare WF, HU Ishoff POTHE, Haag J, Balaz SJ, Dingermans A et al. Differentiating between low and high susceptibility to Schizophrenia in twins: The significance of dermatoglyphic indices in the relation to other determinants of brain development. Schizophr Res. 2001;52:181-93.
- 30. In: Gangane, eds. Human Genetics. 2nd ed. London: Churchill Livingston publications; 1992 and 1996.
- Gapta CM, Tutakne MA, Bhana BA. Absence of triradius 'd' on the palm of leprosy patients. Ind J Leprosy. 1984;56(4):584-91.
- 32. Rajangam S, Janakiram S, Thomas L. Dermatoglyphics in Down's syndrome. J Indian Med Assoc. 1995;93(1):10-3.
- 33. Miglinets V. Relationship between Dermatoglyphic variability and finger length in genetic disorders: Down's syndrome. Genetica. 1991;27(3):541-7.
- 34. Reed T, Reichmann A, Palmer C. Dermatoglyphic differences between 45x and other chromosomal abnormalities of Turner's syndrome. Hum Genet. 1977;36(1):13-23.
- 35. Komotz Y, Yoshida O. Finger patterns and ridge counts of patients with Klinefelter's syndrome (47xxy) among the Japanese, Hum Hered. 1976;26(4):290-7.
- Vishvanathan G, Singh H, Ramanugam F. Dermatoglyphic analysis of fingertip print pattern of blind children from Bangalore. J Ecotoxicol Environ Monit. 2002;12:49-52.
- 37. Rott H, Schwanitz G, Riether M. Dermatoglyphics in Noonan's syndrome (Authors transl). Acta Genet Med Gemellol (Rome). 1975;24(1-2):63-7.
- 38. Weinre H. Finger print pattern in Alzheimer's disease. Arch Neurol. 1995;42(1):50-4.
- 39. Pahuja K, Agarwal SK. Analysis of quantitative and qualitative dermatoglyphic traits in Schizopherinic patients. J Anat Soc of India. 2012;61(2):269-72.
- Simsek S, Taskiran H, Karakaya N. Dermatoglyphic analysis in children with CP. Neurobiology – BP. 1998;6(3):373-80.
- Pallotta R, Carlone G, Petrucci A, Chiarelli F. Dermatoglyphics in Von Recklinghausen neurofibromatosis. Amer J Med Genetics. 1989;34(2):233-6.
- 42. Lal N, Surekha RK. A study of dermatoglyphic pattern in epileptic patients. J Anat Soc India. 2012;61(1):26-9.
- 43. Nair R. Dermatoglyphic diversity in congenital heart diseases. Indian J Medical Res. 1986;83:56-67.
- 44. David TJ. Dermatoglyphics in congenital heart disease. J Med Genet. 1981;18(5);344-9.
- 45. Annupurna V, Ahuja YR, Reddy GD, Rao VS, Rao PN. Dermatoglyphics studies in Rheumatic heart disease. Hum Hered. 1978;28:72-8.

- 46. Sant SM, Vare AM, Fakhruddin S. Dermatoglyphic traits in Diabetes mellitus. J Anat Soc India. 1980;29:43.
- 47. Ravendranath R, Thomas IM. Finger ridge count and finger print pattern in maturity onset diabetes mellitus. Ind J Med Sci. 1995;49:153-6.
- 48. Pal GP, Roufal RV, Bhagvat SS. Dermatoglyphics in carcinoma cervix..J Ant Soc India. 1985;34(3):157-61.
- 49. Gupta CM, Tutakne MA. An evaluation of palmer flexion creases and dermatoglyphics in Leprosy. Indian J Lepr. 1986;58:263-75.
- 50. Pursani MZ, Elhence GP, Tibrewala L. Palmer dermatoglyphics in essential hypertension. Indian heart J. 1989;41:119-22.
- 51. Gupta UK, Prakash S. Dermatoglyphics: a study of fingertip patterns in Bronchial asthma and its genetic disposition. Kathmando Univ Med J. 2003;1(4):267-71.

- 52. Ravindranath R, Shubha R, Nagesh HV. Dermatoglyphics in Rheumatoid arthritis. Ind J Med Sci. 2003;57:437-41.
- 53. Babu SS, Powar BP, Khare ON. Palmer dermatoglyphics in pulmonary tuberculosis. J Ant Soc India. 2005;54(2):1-9.
- 54. Chintamani, Khandelwal R, Mittal A, Saijanani S, Tuteja A, Bhansal A, Bhatnagar D, Saxena S. Qualitative and quantitative dermatoglyphic traits in patients with breast cancer. A prospective clinical study. BMC Cancer. 2007;7:44.
- 55. Oladipo GS, Olabiyi O, Oremosu AA, Noronha CC, Okanlawon AO, Paul CU. Sickle cell anaemia in Nigeria: dermatoglyphic analysis of 90 cases. African journal of Biochemistry Research. 2007;1(4):54-9.

DOI: 10.5455/2320-6012.ijrms20140207 **Cite this article as:** Bhat GM, Mukhdoomi MA, Shah BA, Ittoo MS. Molecular dermatoglyphics: in health and disease - a review. Int J Res Med Sci 2014;2:31-7.