

Review Article:

Loss of fingerprints: A hidden problem facing during biometric identification

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Abstract

In the era of globalization as well as digitalization fingerprinting is the most acceptable form of digital recognition. It is the most available unique method for identification and individuality. But in recent years people from different corners of the world have noticed to have no fingerprinting incidentally when they are facing problems in digital identification and authentication systems. A thorough literature review was conducted involving different medical databases, using the following terms: loss of fingerprints, adermatoglyphia, and biometrics both individually and in combination. Surprisingly, no articles were found discussing specific guidelines for adermatoglyphic patients who have to undergo mandatory fingerprint verification. However, few case reports and scientific studies reveal that it will be a burning issue within the next decade. This article highlighted the epidemiology, types of adermatoglyphia, and challenges in diagnosing the cases and the importance of imposing other biometric methods to overcome the problem.

Keywords: Adermatoglyphia, fingerprints,

Introduction

The scientific and methodical study of complex patterns and fingerprints from palms, fingers, soles and toes is referred to as dermatoglyphics.^{1,2,3} Dermatoglyphics (or fingerprints) is a Greek word meaning skin (derma) and carvings (glyphe). These ridges are distinctive carvings of nature especially prominent over the palmar surface of hands and plantar surface of feet.⁴ More than a century ago, Sir Francis Galton discovered that these ridge patterns are incredibly constant throughout the lifespan of an individual.⁵ Later, the term "dermatoglyphics" was introduced by Dr. Harold Cummins in 1936.⁶ All over the world, fingerprinting is the most widely used method for individual identification and authentication (I&A). Dermatoglyphics form a major portion of mass data collection, biometric surveillance, and verification. Adermatoglyphia is defined as the congenital or acquired loss of the epidermal ridge pattern.⁴ It can be complete or partial, reversible or irreversible. It is also referred to

as "Immigration Delay Disease".⁷ Adermatoglyphia, or simply, loss of fingerprints attributed to a medical cause, represents a taxing situation for such biometric scrutiny systems requiring a fingerprint scan as a mandatory phase in identification and authentication procedure. The scenario can be extremely debilitating for adermatoglyphic patients, especially when the condition is permanent or irreversible.⁸

The disparity in ridge pattern of every individual is partly determined by genetics during gestational weeks 7th to 21st and partly by changes, trauma and diseases acquired during life.^{9,10} Biometric methods which are used for identification in recent days are complex, however, the use of fingerprints is less complex, cost-effective, readily available and therefore, universally used compared to other modalities. However, numerous hereditary and acquired causes of adermatoglyphia pose a huge

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hurdle in individual identification.¹¹

Epidemiology

Adermatoglyphia is not a known topic and no extensive work has not been done on it. So, there is a dearth of literature. However, nowadays the need for biometrics in identification, authentication and forensics application have exposed cases of hidden adermatoglyphia and the field of interest is increasing. Haber et al. conducted a national survey in Lebanon in collaboration with the Ministry of internal affairs in 2014 which recorded 0.18%

cases of adermatoglyphia with a female predominance and an increased incidence in the ageing population with dermatitis as the commonest cause.¹²

A cross-sectional survey was carried out in the Department of Dermatology, Services Hospital, Lahore. All patients were referred by the National Database and Registration Authority (NADRA) through the Medical Examination Department (MED) of Services Hospital for verification of adermatoglyphia. A total of 152 patients with adermatoglyphia were enrolled over two years. Out of them, 96 (63.2%) were males while 56 (36.8%) were females. The mean age of the patients was 41.89 ± 23.72 years. Among their study population, 20 (13.2%) patients had isolated congenital (idiopathic) adermatoglyphia, 52 (34.2%) patients suffered from Congenital adermatoglyphia associated with a syndrome, while 80 (52.6%) patients presented with acquired causes of adermatoglyphia.¹¹

Loss of fingerprints can be congenital or acquired.^{8,13} The congenital form can be part of a complex syndrome and the prevalence will mirror that of the syndrome. Individuals with completely missing fingerprints as an isolated finding is extremely rare. Four generations were reported to have this isolated form of adermatoglyphia inherited in an autosomal dominant fashion.¹⁴

The acquired forms are more common and could result from other dermatological conditions.^{8,14-15}

A 27-year service holder visited in a private chamber referred by a travel agency in July 2023 for medical evaluation. He realized the loss of his fingerprints when he was unable to give his fingerprint during passport issuing, requiring his to be ten-printed (i.e., all fingers and thumbs of both hands were scanned/printed). Physical examination reveals hyperhidrosis of both palms and laboratory reports reveal no abnormalities. He was treated as a case of idiopathic palmar hyperhidrosis and advised to follow up after 2 months.



Loss of fingerprints of a 27 years male due to palmar hyperhidrosis

BBC reported in December 2020, that a Bangladeshi family where 3 generations had no fingerprints faced obstacles in issuing national ID cards, driving licences, buying SIM cards and office attendance systems. A dermatologist in Bangladesh has diagnosed the family's condition as congenital palmoplantar keratoderma. More genetic testing would be needed but these were not available in this resource-poor country. They got their NID card issued by the Bangladesh Government after presenting a medical certificate. The card used other biometric data too - retina scan and facial recognition.¹⁶



Congenital palmoplantar keratoderma of a Bangladeshi male¹⁶

The rationale of the review

Clinically adermatoglyphia is defined as the absence of an epidermal ridge pattern that may be due to congenital or acquired causes.⁶ A few or all digits can be affected. It may also refer to the absence of the ridge patterns formed on the plantar aspects of the feet. Furthermore, adermatoglyphia can refer to a partial loss of the ridges (i.e., ridges are unnoticeable on general evaluation but detectable on deep inspection or under a magnifying lamp) or a complete absence (depicting total effacement) of epidermal ridges. These are time-consuming challenges not only for the concerned authorities but also for the individual who has to face the problem in completing verification procedures. Face recognition and fingerprinting are the primary modes of biometric I&A

all over the world. Any technical problem or error hindering these steps of verification can cause the entire verification process to come to a halt.¹² Most patients with adermatoglyphia are unaware of the fact that they cannot produce their fingerprints and are surprisingly shocked when biometric analysis fails. Nothing can be done except resorting to other modalities of identification. This is also a challenge in other areas such as forensics and criminal identification as it slows the process. Diagnosing adermatoglyphia is quite challenging. Therefore, a reliable alternative to fingerprinting that is cheap, unique and readily available must be sought shortly.

Table 1. Congenital causes of adermatoglyphia and their associated genes¹⁷

S/no	Disease condition	Associated Gene
1	Basans syndrome	Smarcad1
2	Naegeli-franceschetti-jadassohn syndrome	Krt14
3	Dermatopathia pigmentosa reticularis	Krt14
4	Reticulate acropigmentation of Kitamura	Krt14, dkc1
5	Rothmund thomas syndrome	Wrn,blm
6	Dyschromatosis universalis hereditaria	Smarcad1,dkc1

Table 2. Dermatological and non-dermatological causes of acquired adermatoglyphia¹⁷

Dermatological	Non dermatological
Allergic and irritant contact dermatitis	Trauma
Atopic dermatitis	Burn
Dyshidrotic eczema	Amputation
Cutaneous LE	Caustic abrasion
Epidermolysis bullosa	Denervation injuries
Pemphigus vulgaris	Capecitabine
Psoriasis	Topical steroid
Keratoderma blennorrhagica	Retapamulin
Palmar wart	Atorvastatin
Leprosy	
Pyoderma/impetigo	
Coxsackie	
Tinea mannum	
Erythema multiforme	
Steven johnson syndrome	
Toxic epidermal necrolysis	
Serum sickness	
Primary hyperhidrosis	
Lichen	
Acrodermatitis	

Methods of human identification and authentication

The necessity of a second-line substitute I&A system, especially for patients suffering from irreversible adermatoglyphia can only be realized when people get into the hassle of identification. Imperatively a medical certificate can be medico-legally issued but only provides a temporary solution, as the validity and the authenticity of such certificates might not be acceptable by other organizations. Different set-ups demand different types of biometric verification, often involving mandatory, multi-modal biometric functions operating without a backup plan.^{18,19} Additionally, some corporations require only thumb-printing for monitoring office attendance or bank account verification. Other agencies granting citizenship, driver’s licenses, passports, and immigration papers require a more stringent policy, requiring individuals to be ten-printed. Therefore, a substitute I&A system is recommended that can function globally as a default program for patients suffering from irreversible adermatoglyphia.

Biometrics

Over the past few decades, many advancements have been made in biometric technology with the introduction of new measurable biological traits as potential biometric targets. Table 3 presents a concise review of the indispensable characteristics required for a biometric indicator.

While many of these biological traits fulfil the necessary characteristics required to qualify as a biometric indicator, feasibility becomes the most important factor based on efficacy, social acceptability, the technical complexity of systems, ease of applicability, and cost-effectiveness. Consequently, there is no single biometric indicator that ideally achieves all these parameters. Therefore, hybrid or complex multi-modal biometric systems are used to enhance the accuracy and efficacy of I&A.^{18,19}

TABLE 3: Essential qualities of a biometric indicator

Characteristic	Definition
Generality	Universally present in all individuals
Uniqueness	No individuals share the same configuration
Stability	Unchanging throughout the lifespan
Quantifiable	Measurable for comparison

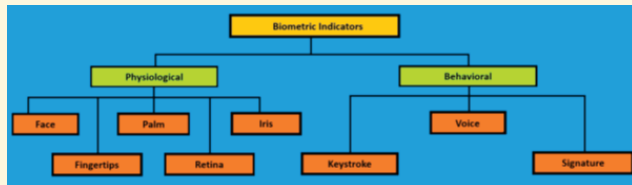


Fig: Biometric Subtypes

Classification is based on physiological and behavioural traits

In addition to facial recognition and fingerprinting technology, extensive research has been conducted on other biometric recognition techniques, namely, iris/retinal scanning, hand geometry and palm prints, DNA fingerprinting, voice recognition, signature scanning, gait

patterns, and keystroke patterns.²⁰⁻²³ Future endeavours will involve the exploration of other anatomical organs and parts, with the potential for I&A to fulfil essential biometric features. These domains include but are not limited to, finger-vein patterns,²¹ foot-printing,²⁴ dorsal finger pattern,²⁵ lip-printing,²⁶ and tongue-printing.²⁷

Challenges

Diagnosing adermatoglyphia in Bangladesh is quite challenging. Patients usually present with difficulty in capturing fingerprints during biometrics. Further evaluation to enable the Attending physicians cannot get a chance to find out a specific cause because of the lack of molecular genetic testing. This is the primary investigation to ascertain whether the patient has smarcad1 gene and keratin¹⁴ mutation which are the major gene mutations seen in adermatoglyphia.^{28,29} These investigations are not readily available and definitive diagnosis is halted. Other investigative modalities like volar pad biopsy are usually refused by patients as they see the absence of fingerprints as a variant of normal and will resist attempts for invasive procedures. Another challenge is tracing family members to ascertain whether they also have adermatoglyphia.

The paucity of data on adermatoglyphia is also an issue as most patients with the condition do not present to a health care facility except when in need of a medical report. Patients are usually seen during biometrics in the banks, borders or immigration offices and access to these data is met with some bureaucracies. The psychosocial impact patients with adermatoglyphia face during biometric screening is also enough

trauma to make them avoid further analysis.

Conclusion

Adermatoglyphia, especially due to acquired causes is not an uncommon finding in the geriatric age group and manual labourers. It is also found in the young population. Despite its benign nature, the persistent invasion of biometric identification in modern life via fingerprint verification creates huge hassles for patients with irreversible adermatoglyphia. As fingerprinting is the most available and cheaper method for biometric identification, it provides the identification and authentication system quicker and is applied in different systems and institutions. However, a person with absent or permanent loss of fingerprinting has to undergo enormous hardships due to lack of any alternative options. Therefore, specific guidelines or a substitute I&A system is recommended that can function worldwide, as a default program, facilitating I&A for patients suffering from irreversible adermatoglyphia.

Conflict of Interest

None

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Patient Consent

Taken

IRB approval status

Not applicable

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