Understanding the formidable nail barrier: A review of the nail microstructure, composition and diseases

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Summary
The topical treatment of nail fungal infections has been a focal point of nail research in the past few decades as it offers a much safer and focused alternative to conventional oral therapy. Although the current focus remains on exploring the ways of enhancing permeation through the formidable nail barrier, the understanding of the nail microstructure and composition is far from complete. This article reviews our current understanding of the nail microstructure, composition and diseases. A few of the parameters affecting the nail permeability and potential causes of the recurrence of fungal nail infection are also discussed.

KEYWORDS
anti-fungal, iontophoresis, keratin, laser treatment, nails, onychomycosis

1 | INTRODUCTION

The nail is modified to a great extent in some mammals, for example, as a prehensile organ—the claw, or as a point of locomotion—the hoof. In humans, the nail plate is the completely keratinised part of the upper surface of the tip of each finger and toe which provides protection to the phalanges, enhances dexterity and facilitates scratching. In addition, nails serve an aesthetic and cosmetic purpose.

The development of nail unit in humans takes place in between the 9th and 17th weeks in utero by an intricate series of mesenchymal-ectodermal interactions.1 The individual digits are apparent from the 8th week of gestation.2 Nail anlage, the epidermis overlying the dorsal tip of the digit, presents from 9 weeks and is the first embryonic element of the nail unit. A distinct region called the primary nail field can be seen at 10 weeks overlying the tip of terminal phalanx. At the 13th week, proximal and lateral nail folds develop as a result of differential slow growth of the primary nail field and surrounding tissue. By the 14th week, the nail plate can be seen emerging from under the proximal nail fold. At the 17th week, most of the nail bed is covered by nail plate. From week 20, finger and nail grow approximately in tandem.

The growth of mature adult nails has been studied extensively.3,4 Normal fingernail growth varies from less than 1.8 mm to more than 4.5 mm/mo and varies distinctly between individuals, but is more consistent among members of the same family.5 The average nail growth rate is 0.1 mm/d (3 mm/mo) and is used to predict nail regrowth time.6 A normal fingernail grows out completely in about 6 months, whereas toenails grow one-third to half the rate of fingernails; thus, toenails take 12-18 months to grow out entirely.7 The nail growth rate is less than normal in people who are immuno-compromised, immobilised or paralysed, malnourished, suffering from acute infection or undergoing anti-mitotic drug therapy.8

2 | STRUCTURE OF HUMAN NAIL UNIT

Figure 1 shows a schematic diagram of human nail unit which is comprised of four main structures: the proximal nail fold (PNF), nail matrix, nail bed and hyponychium. These structures together form nail plate—the flat, rectangular, translucent and keratinised structure sitting on the upper surface of the tip of each finger and toe. The cells in the nail plate are closely packed, adherent, interdigitating and very flat, lying perpendicular to the plane of the nail plate surface.

2.1 | The proximal nail fold (PNF)

The PNF is a wedge shaped fold of skin lying on the dorsum of finger and toes which covers approximately one-fourth of the total nail length. The nail plate appears to emerge out of the PNF. The skin on
either side of the nail is an extension of the PNF, and is known as the lateral nail fold. It comprises two layers of epidermis; the dorsal layer which forms the dorsum of finger epidermis and the ventral portion which forms eponychium. The stratum corneum of eponychium forms a thick rim of keratinous material, known as cuticle, around the margins of the PNF which along with the PNF and lateral folds provide protection against penetration of water and toxic chemicals. The disruption of cuticle can lead to bacterial or fungal infection resulting in a nail disease called paronychia, which is characterised by the inflammation of the PNF.

### 2.2 The nail matrix and nail plate

Directly underneath the PNF is a small area of highly proliferative epidermal tissue, known as the nail matrix, which produces the nail plate. Nail plate is made up of three different layers which are well-bonded together. The visible white area of nail matrix is called lunula. The nail plate pigmentation varies with race and is defined by melanocytes present in the nail matrix. Langerhan’s cells are also present in the nail matrix. Nail Matrix produces the nail plate which comprises of an average of 196 cell layers that are distributed among three tightly bound layers—dorsal, intermediate and ventral (Figure 2). The thickness ratio of the three layers of nail plate was found to be 3:5:2 by Kobayashi et al. In the same study, dorsal plate was stated to be the rate limiting barrier for permeation of 5-fluorouracil (5-FU), when compared with intermediate and ventral layers. This observation has not been independently confirmed, either with 5-FU or with other compounds. Examination of the biochemistry of nail plate suggests that the interpretation of dorsal layer being the primary barrier for nail permeation should be viewed cautiously. Although the sulphur concentration is equivalent in the dorsal and intermediate plates, sulphhydril groups are predominant in dorsal plate, whereas the amino acid cystine, which contains disulphide bonds, is concentrated in the intermediate plate. This indicates a higher degree of cross-linking in the intermediate plate, which limits swelling and suggests that its barrier properties are likely to be significant.

Of the three main classes of protein (fibrous, globular and conjugated), the nail plate is composed of keratin, a scleroprotein (fibrous protein) which contains large amounts of sulphur. Polypeptide chains are the basic macromolecules which form keratin. These chains exist either in curled helical conformation (α-keratin) or side-by-side pleated conformation (β-keratin). In mammals, α-keratins are the primary constituents of hair, nails, hooves, quills, horns and the epidermal layer of the skin. Two distinct types of α-keratin exist, (a) hard α-keratins, derived embryologically from epithelial cells, are the main constituents of cornified mammalian tissues such as nails, hooves, horns, quills, and (b) soft α-keratin, is the main component of epidermis (of skin) and other epithelial tissues. Human nails are comprised of approximately 80% hard α-keratin and 20% soft α-keratin. The dorsal plate cells have thickened plasma membranes and are anucleated as shown by the lack of acid phosphatase activity. The intermediate plate possesses some eosinophilic cytoplasm and nuclear remnants. They are held firmly together by desmosome glue spots and complex interdigitations. The cells of the ventral plate emerge from the nail bed and are an easy target for disease.

The keratin intermediate fibres/filaments (~70 Å diameter) in the intermediate layer are well aligned and oriented in the transverse direction laterally, perpendicular to the axis of nail growth, whereas keratin fibres in the dorsal and ventral plates are packed like overlapping tiles and have no preferred orientation (Figure 3). The toughness of the nail plate can be attributed to this alignment of keratin fibres and to the presence of large numbers of disulphide cross-links due to high proportions of cystine.

The microstructure of nail plate was further analysed in detail by observing effects of keratinase enzyme on nail plate. It was observed that the enzyme acted on the intercellular matrix that holds the nail corneocytes together and the nail corneocytes on the dorsal...
surface appeared to be separated from each other. In contrast, the nail corneocytes were not differentiated in control nail sample immersed in buffer solution as they were closely bound to each other. The deeper layers of nail plate (intermediate and ventral) were not affected by the enzymatic action probably due to hindered diffusion of large molecular size (MW~33,000) of the enzyme to these layers.

To supplement the microstructural findings of Mohorcic et al., SEM imaging on nail plate was also performed in collaboration with Forensic Science Lab at Amway Corporation, Ada, MI (Figures 4, 5, and 6). SEM imaging was performed on nail clipping under three different conditions—(i) room temperature—Figures 4A-F and 5A; (ii) hydrated nail after 24 hours—Figure 5B; and nail treated with 5% w/w Solvable® tissue dissolving fluid (after 4 hours—Figure 6A-B). The morphological features of the three layers were found to be similar as mentioned earlier.

The surface appearance of dry nail clipping at room temperature was found to be different than hydrated nail clipping. The dry nail clipping showed stratified corneocyte surface with ridges on the dorsal side, probably representing bundles of keratin fibres (Figure 5A). In comparison, the hydrated nail plate showed less striation/ridges on the dorsal surface (Figure 5B).

Based on the autoradiographic data, the thickness of the nail plate is determined by the length of the matrix. The thickness of the nail plate progressively increases from proximal to free distal end.
The average nail plate thickness is 0.5 mm in women and 0.6 mm in men. The nail plate thickness progressively increases with age as a result of increasing size of the cells in the plate and decreased growth rate.\(^5\)

### 2.3 The nail bed

The nail bed is a pink-colored soft, thin, noncornified epithelium surface, devoid of granular layer and sebaceous glands, which lies below the nail plate and extends from the nail matrix to the hyponychium. Sometimes, it is referred to as the ventral matrix, since it contributes a few layers of cells on the ventral surface of the nail plate. It is considered to be a transitional zone where living cells keratinize and are integrated in the nail plate. Also, the nail bed is known to have corpuscles and nerve endings which are responsible for sensitivity to light, touch, and pressure.\(^9\) The pink color of the nail bed is due to the enriched vascular supply just below the nail bed.

### 2.4 The hyponychium

The hyponychium is the thickened epidermis beneath the free distal end of the nail of a digit. It performs the same protective function as provided by the cuticle, PNF and lateral folds.

### 3 FORMATION OF NAIL PLATE

The nail matrix constantly replenishes the nail plate. The nail matrix is comprised of stratified squamous epithelial cells which are held together by long rete ridges and sparse dermis. Mitosis in the basal layer of the nail matrix constantly replaces the matrix. The matrix cells differentiate and move continuously, and consequently flatten by dumping their organelles and condensing their cytoplasm. The pressure of this new buildup forces the other cells (nail plate) to move dorsally (Figure 7). In due course, the cells keratinise as the nuclei vanish.\(^22\) Cell division of nail matrix is continuous; thus, nail is formed continually, unlike hair which is formed cyclically. The cells at the dorsal surface of nail plate overlap to form a smooth surface, in contrast to the ventral surface which is irregular allowing interdigitation with the nail bed. The matrix turnover rate defines the growth rate of nail. The nail form is determined primarily by the growth rate of the dorsal, intermediate and ventral matrices, and could be affected by various disease conditions (Figure 8).

### 4 BIOCHEMISTRY OF THE HUMAN NAIL PLATE

The keratins found in human nail plate are almost identical to those of hair,\(^23,24\) which is primarily composed of \(\alpha\)-keratin. The analysis of human nail plate by immunoblotting technique has shown that it is comprised of approximately 10%-20% soft epithelial keratins (50K/58K and 48K/56K keratin pairs) with the remainder being hard \(\alpha\)-keratin.\(^15\) Despite the existence of different forms of \(\alpha\)-keratins, they exhibit many similarities in their chemical and structural features. Figure 9 demonstrates the basic molecular structure of \(\alpha\)-keratin. The isolated \(\alpha\) helical chains (formed by condensation of amino acids) form a coiled coil dimer with cross-linked disulphide bonds to assemble into a protofibril unit. Eight protofibril units polymerise together to form the basic structural unit, the intermediate filament (IF) or microfibril. In human hair, \(\alpha\)-keratins have unique arrangement of aligned
IFs (~7 nm diameter and spacing of 10 nm) and are enclosed by a non-filamentous protein matrix (Figure 10). These hair-like intermediate filaments which exist only in the intermediate layer of the nail plate are measured to be approximately 7 nm in diameter by various researchers.14,27 The polypeptide chains in keratin are formed by condensation of several amino acids. Although all are derived from the ectodermal cells, fundamental physical and chemical differences in fully differentiated forms of nail, hair and stratum corneum exist. A comparison of amino acid composition of human nail, hair and stratum corneum is shown in Table 1.19,28 Hair and nail are differentiated along the same lines despite their different morphological characteristics.9 Nail contains 10%-20% more basic soft keratins, in addition to the sulphur-rich hard keratins present in both tissues.15 A significant proportion of glutamic acid, half-cystine, arginine, aspartic acid, serine and leucine are present in the human nail plate. It is noteworthy to observe that the content of glycine and half-cystine is similar in human hair and nail, but not stratum corneum. An outstanding degree of physical and chemical stability is conferred to the keratin filaments by the extensively cross-linked disulphide structures corresponding to the high proportions of cystine.19

The proteins of nail and hair are alike and have extensive folding supported by extremely stable disulphide bonds and in addition, have Van der Waals interactions, hydrogen bonds and coulombic interactions. This explains the high degree of chemical and physical resistance of nail and hair, in contrast to the skin. Although these disulphide bonds are also found to exist in skin to a lesser extent, the geometric spatial configuration of these bonds is different for nail/hair and skin. As determined by Raman spectroscopy, hair and nail have these bonds in gauche-gauche-gauche conformation and those for stratum corneum are in gauche-gauche-trans conformation, with the latter being less stable.29

The extreme folding of protein molecules and different geometries of disulphide bonds in nail/hair and skin has resulted in a different degree of hydration. Water has been stated to exist in the nail plate in either free or bound form, with the latter accounting for most of the nail's water content.29 In comparison with the normal nail structure, the loose skin structure allows more free water than bound water. More quantitative analysis indicates that very little of the water in the nail plate can be considered to be “free”.30 The uptake or rate of loss of water is not affected by the extraction of lipids which suggests that the role played by the lipid pathway in diffusion process of water across the nail plate is insignificant.31 Although the detailed characterisation of nail keratins has not been done to date, one can approximate a weighted average isoelectric point (pI) of the nail plate by using the per cent composition and the individual pI's of the amino acids from Table 1. This value was calculated to be 4.7.

The amounts of minerals and electrolytes found in nail plate are highly variable. They include elements such as sodium, potassium, nitrogen, magnesium, calcium, iron, copper, zinc, phosphorus, selenium, sulphur and several others.9,10,22 Table 2 gives an overview of different methods employed to determine minerals and other exogenous materials in the nail plate. Cholesterol is the primary lipid component of the nail plate. In contrast to the 10% lipid content and 20%-40% water content in the stratum corneum, nail plate contains 0.1%-1% lipids and 7%-12% water (varies with relative humidity).9

**FIGURE 9** Molecular structure of α-keratin. (i) A dimeric coiled coil formed by two keratin polypeptides; (ii) tail-to-head association of coiled coils to form protofibril and (iii) eight protofibrils associate together to form an intermediate filament (IF)

**FIGURE 10** Schematic diagram of a human hair fibre
**Table 2** Different methods for constituent analysis of nail plate

<table>
<thead>
<tr>
<th>Method</th>
<th>Element</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>(I) Structural and mineral constituents</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Raman spectroscopy</td>
<td>Water, protein and lipid</td>
<td>29</td>
</tr>
<tr>
<td>Immunohistochemistry</td>
<td>Keratin</td>
<td>100</td>
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<tr>
<td>Polymerase chain reaction</td>
<td>Deoxyribonucleic acid</td>
<td>101,102</td>
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<tr>
<td>Electron microscopy</td>
<td>Cystine</td>
<td>103</td>
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<tr>
<td>X-ray diffraction</td>
<td>Mg, Cl, Na, Ca, S, Cu</td>
<td>34,104</td>
</tr>
<tr>
<td>Colorimetry</td>
<td>Fe</td>
<td>105</td>
</tr>
<tr>
<td>(II) Exogenous materials</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Atomic absorption spectroscopy</td>
<td>Cd, Pb, Zn, Ca, Cr, Fe, Cu, Cr, Fe, Mn, Ni, Co, Na, K</td>
<td>34,106,107</td>
</tr>
<tr>
<td>Mass fragmentography</td>
<td>Metamphetamine</td>
<td>108</td>
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<tr>
<td>Gas chromatography</td>
<td>Amphetamine, Cocaine</td>
<td>109,110</td>
</tr>
<tr>
<td>Flow injection hydride generation atomic absorption spectrometry</td>
<td>Arsenic</td>
<td>111</td>
</tr>
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</table>

**6 | Electrical Properties of Human Nail Plate**

In vivo and in vitro resistance of nail plate immersed in phosphate-buffered saline (PBS) solutions as a function of time (hydration) was studied by Hao et al. The resistance of nail plate in both in vivo and in vitro experiments decreased significantly upon 2-hour nail hydration and then slowly decreased to a constant value over time. A significant decrease in nail resistance with hydration supports the notion that nail hydration plays a significant role in the diffusion process through the nail.

**7 | Permeability of Human Nail Plate**

Differences in the physical and chemical characteristics of nail and stratum corneum translate to differences in solute permeation through the membrane. Although the nail plate has been historically considered as a fibre matrix system rather than a lipophilic membrane, a recent report by Baswan et al. suggested that there might be additional structure to the nail plate such as residual lipids which leads to lower permeability of hydrophilic to moderately lipophilic solutes. The diffusion of hydrophilic solutes may be partially impeded by the remnants of the cell membranes which may form a discontinuous lipid barrier. Permeation of drugs through the nail plate depends on various parameters such as the physico-chemical properties of the drug, nail properties such as hydration and thickness, the nature of the

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**Table 1** Amino acid composition (as residues per 100 residues) of human nail, hair and stratum corneum

<table>
<thead>
<tr>
<th>Amino acid</th>
<th>pI</th>
<th>Nail (%)</th>
<th>Hair (%)</th>
<th>Stratum corneum (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lysine</td>
<td>9.74</td>
<td>3.1</td>
<td>2.5</td>
<td>4.2</td>
</tr>
<tr>
<td>Histidine</td>
<td>7.59</td>
<td>1.0</td>
<td>0.9</td>
<td>1.5</td>
</tr>
<tr>
<td>Arginine</td>
<td>10.76</td>
<td>6.4</td>
<td>6.5</td>
<td>3.8</td>
</tr>
<tr>
<td>Aspartic acid</td>
<td>2.77</td>
<td>5.4</td>
<td>7.9</td>
<td>7.9</td>
</tr>
<tr>
<td>Threonine</td>
<td>5.60</td>
<td>6.1</td>
<td>7.6</td>
<td>3.0</td>
</tr>
<tr>
<td>Serine</td>
<td>5.68</td>
<td>11.3</td>
<td>12.2</td>
<td>13.6</td>
</tr>
<tr>
<td>Glutamic acid</td>
<td>3.22</td>
<td>13.6</td>
<td>12.2</td>
<td>12.6</td>
</tr>
<tr>
<td>Proline</td>
<td>6.30</td>
<td>5.9</td>
<td>8.4</td>
<td>3.0</td>
</tr>
<tr>
<td>Glycine</td>
<td>5.97</td>
<td>7.9</td>
<td>5.8</td>
<td>24.5</td>
</tr>
<tr>
<td>Alanine</td>
<td>6.00</td>
<td>5.5</td>
<td>4.3</td>
<td>4.4</td>
</tr>
<tr>
<td>Valine</td>
<td>5.96</td>
<td>4.2</td>
<td>5.5</td>
<td>3.0</td>
</tr>
<tr>
<td>Methionine</td>
<td>5.74</td>
<td>0.7</td>
<td>0.5</td>
<td>1.1</td>
</tr>
<tr>
<td>Isoleucine</td>
<td>6.02</td>
<td>2.7</td>
<td>2.3</td>
<td>2.7</td>
</tr>
<tr>
<td>Leucine</td>
<td>5.98</td>
<td>8.3</td>
<td>6.1</td>
<td>6.9</td>
</tr>
<tr>
<td>Tyrosine</td>
<td>5.66</td>
<td>3.2</td>
<td>2.2</td>
<td>3.4</td>
</tr>
<tr>
<td>Phenylalanine</td>
<td>5.48</td>
<td>2.5</td>
<td>1.7</td>
<td>3.2</td>
</tr>
<tr>
<td>Half-cystine</td>
<td>5.07</td>
<td>10.6</td>
<td>15.9</td>
<td>1.2</td>
</tr>
</tbody>
</table>

*pI is the isoelectric point of the individual amino acids.*

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Forsslind that the effective elastic modulus of nail is directly related to cell arrangement, cell adhesion, microstructure of keratin fibres and hydration level. Farren et al. showed that cutting nail longitudinally requires double the energy (~6 kJ/m²) than cutting it transversely (~3 kJ/m²) and that the upper layers of nail plate provide most of the nail resistance against tearing.
formulation (eg, pH and use of chemical penetration enhancers) and the binding of drug to keratin matrix, which have been widely studied by various researchers.12,35-37,40-50

7.1 | Effect of lipophilicity and molecular weight of the permeant

Walters et al.26,37 performed transungual permeation studies for a homologous series of alcohol ranging from methanol to dodecanol and compared it to stratum corneum data. As the alkyl chain length increased from 1 to 8, the permeability coefficient through the nail plate decreased. This result was attributed to a decrease in the partitioning of solutes in the nail plate. They also suggested the existence of a parallel lipid pathway for extremely lipophilic compounds, since the permeability coefficient of alkanols after n-octanol increased with chain length. Studies conducted by Kobayashi et al.52 suggested that the dependence of permeability on molecular weight is more significant than lipophilicity. This conclusion was also supported by a more comprehensive dataset of 42 compounds analysed by Baswan et al.38,39. Studies conducted by Mertin and Lippold45 on 10 anti-fungal drugs also showed that hoof permeability decreased with an increase in molecular weight of the drug. The existence of a parallel lipid pathway as proposed by Walters et al. was not supported by studies conducted by Kobayashi et al.

7.2 | Effect of pH

The effects of pH on nail permeability using ionisable drugs have been studied by various researchers. The ionisation of drug has been associated with a decrease in transungual permeability due to an apparent increase in the hydrated radius of the molecule.43,45 Studies conducted by Soong47 in nail plate demonstrated that the permeability coefficient of benzoic acid (a weak acid) decreased by 95.5% as pH was increased from 2.0 to 8.5, indicating that maximum permeation rate was achieved for undissociated benzoic acid at a lower pH. Similar studies were conducted by Mertin and Lippold45 in bovine hoof membrane using benzoic acid and pyridine (a weak base) at pH 2.0 and 7.4. The authors attributed the enhanced permeability of benzoic acid at pH 2.0 and pyridine at pH 7.4 to Donnan equilibrium effect of nail plate. As the isoelectric point of the keratin is approximately 5, it is negatively charged at pH 7.4 and positively charged at pH 2.0. Thus, it hinders transport of positively charged pyridine molecule at pH 2.0 and enhances the permeation of undisassociated benzoic acid and vice-versa at pH 7.4. Kobayashi et al.43 demonstrated that the dissociation of the drug leads to reduction of permeability in nail plate, irrespective of the charge on the drug. A magnitude of difference on log scale in the regression lines (permeability vs molecular weight) of ionic and non-ionic drug was observed. On the contrary, Walters et al.36,49 studied the nail permeation of anti-fungal drug miconazole and concluded that both ionised and neutral form penetrated the nail at comparable rates.

Similar to passive diffusion, transungual iontophoresis is also affected by the change in pH. The nail plate exhibited iontophoretic permeability for the delivery of glucose and griseofulvin, similar to that observed in human skin.44,52 The permselectivity of nail plate is attributed to the nature of keratin protein whose isoelectric point is around pH 5 and thus, the nail is net negatively charged at pH above the isoelectric point. The anodal iontophoresis was favoured at pH>5 for both glucose and griseofulvin, as the nail plate is negatively charged and enhanced the transport of glucose by convective water flow from anode to cathode direction. It is also noteworthy to observe that passive flux of griseofulvin did not vary significantly with pH. The extent of keratin binding with griseofulvin (33±3.1%) was found to be independent of pH.

7.3 | Effect of nail hydration

The nail plate hydration plays a significant role in permeability of the nail plate. Water flux in fully hydrated nail plate was found to be 12.6±5.6 mg/cm²/h or 126±56 g/m²/h (permeability coefficient of (16.5±5.9)×10⁻³ cm/h) by Walters et al.,48 which is almost 6.5 times the value of 2.5 mg/cm²/h determined by Burch and Windsor in dry nails.53 This is attributed to the fact that hydrated nail plate holds 0.3-0.5 g of water/g of dry nail weight.30,31,41,54,55 Environmental factors such as temperature also influence the permeation in nail plate. Mahothra et al.56,57 found that the water permeability through nail plate is enhanced significantly with an increase in temperature (test conditions: 37, 47 and 57°C). The activation energy of water permeation through nail plate (7.2 kcal/mol) was found to be closer to the value of delipidised human skin (6.3 kcal/mol) than for intact stratum corneum (14.3 kcal/mol). The transonychial water loss (TOWL) was found to be lower for diseased nails (6.9 g/m²/h) than healthy nails (19.4 g/m²/h).58,59 Jemec et al. did not find any correlation between TOWL and nail thickness, whereas Murdan et al.60 and Dutet et al.41 found an inverse linear relationship of TOWL with nail thickness. Also, the values of TOWL in the literature vary from 18-80 g/m²/h.58,60,61 This large variation is attributed to inter- and intra- (different digits) subject variability, environmental conditions and method of measurement. In spite of the difference in TOWL for healthy and diseased nails, Kobayashi et al.53 demonstrated that permeability in nail with fungal infection can be estimated by data on healthy nail plate. Although transepidermal water loss (TEWL) is considered as a marker for epidermal damage, TOWL has not been validated as a marker for nail plate damage.

7.4 | Other enhancement methods

Chemical enhancers such as urea, thioglycolic acid, hydrophobins, 2-mercaptoethanol, N-acetyl-L-cysteine, keratolytic agents and surfactants,41,54,62,63 have shown some promising results in enhancing nail permeability. Other physical methods such as iontophoresis, nail abrasion,69-71 acid etching72 and nail micro-drilling73 also enhance the permeability of the nail plate. Combination of one or more of the above mentioned methods has been reported to yield permeability enhancement synergistically.41,74
The term onychomycosis is derived from the Greek word “onyx” which means nail and “mykes” meaning fungus. The term is used to describe fungal infection of one or more nail units which can be caused by dermatophytes, yeast or non-dermatophyte moulds. The susceptibility of toenail to this disease is 25 times more than the fingernails due to the brunt of the body pressure and occlusive nature of footwear. It is usually asymptomatic and thus patients usually do not have any physical complaints and assume it as a cosmetic problem. As the disease progresses, it can cause pain, discomfort, loss of dexterity and sensation.

8.1 | Aetiology

There are three groups of fungi associated with onychomycosis: dermatophytes, non-dermatophytic moulds and yeast. The rate of involvement of various fungal species in onychomycosis are given in Figure 11. Dermatophytes are the primary causative agents in onychomycosis, accounting for nearly 90% of toenail infections and 50% of fingernail infections. Common fungal agents associated with onychomycosis are:

a. Dermatophytes: *Trichophyton rubrum*, *Trichophyton mentagrophytes* and *Epidermophyton floccosum*


c. Yeast: *Candida albicans*.

8.2 | Clinical presentation of fungal infection in onychomycosis

There are four main clinical presentations of onychomycosis, many of which can be further subdivided into four major types as follows.

8.2.1 | Superficial onychomycosis (SO)

This superficial onychomycosis is primarily caused by dermatophyte *T. mentagrophytes* and sometimes by non-dermatophytic moulds like *Acremonium* spp., *Aspergillus terreus* and *Fusarium oxysporum*. The superficial layers of nail plate are directly invaded by fungi and can be recognised by the presence of opaque yellowish-white islands on the external nail plate. As the viable tissue is not involved in SO, inflammation is minimal in patients.

8.2.2 | Proximal subungual onychomycosis (PSO)

In PSO, the fungal organisms invade the nail unit through proximal nail fold, via cuticle, and penetrate the newly formed nail plate migrating distally resulting in subungual hyperkeratosis, proximal onycholysis and destruction of the nail plate. This fungal infection is visible through the cuticle as yellowish-white discolouration, while the distal nail plate remains normal. It is primarily caused by dermatophyte, *T. rubrum*.

8.2.3 | Distal lateral subungual onychomycosis (DLSO)

DLSO is characterised by the invasion of nail bed and the underside of nail plate by fungi and can be best described as “nail bed dermatophytosis”. The infecting fungi migrate proximally through the underlying nail matrix. It causes mild inflammation and often results in onycholysis and subungal thickening. The common dermatophytes responsible for DLSO are *T. rubrum*, *T. mentagrophytes*, *T. tonsurans* and *E. floccosum*.

8.2.4 | Total dystrophic onychomycosis (TDO)

There are two forms of TDO: primary and secondary. The entire nail unit is involved in TDO and is considered as a combination of all types of the disease.

8.3 | Epidemiology of onychomycosis

The literature available on the epidemiology of onychomycosis is varied. It accounts for approximately 50% of all nail fungal infections and affects 2%-18.5% or more of the world’s population. Age, social class, climate, occupation, predisposing factors, frequency of travel and living conditions determine the prevalence rate of onychomycosis. Studies have shown that prevalence of onychomycosis increases with age [Figure 12].

The reasons for this age-dependent prevalence may be poor peripheral circulation, smoking habits, diabetes, nail trauma, long exposure to pathogenic fungi, suppressed immune functions and inability to maintain good foot care. It is found to be 2-3 times more common in diabetics than non-diabetics. The prevalence of onychomycosis among children varies from 0.44% (United States, Wales and Finland) to 2.6% (Guatemala) and this 30-fold decrease in prevalence rate relative to adults may be attributed to faster
nail growth, small surface area of nail, less time spent in environment containing pathogens and lower prevalence of tinea pedis.  

8.4 | Clinical diagnosis and treatments of onychomycosis

In the present world where advanced medical healthcare is available for most of the conditions, the high prevalence of nail fungal infections suggests that either the treatments currently available are not efficacious or the patients do not realise the condition in early stages. The clinical diagnosis of onychomycosis is based on a physical examination, microscopy culture of nail specimens and PAS (periodic acid-Schiff) examination of subungual debris (highest sensitivity of all methods). The aetiologic agent is identified by culture of sampled nail specimen.  

The treatment of onychomycosis range from palliative to curative therapies which involve the use of oral anti-fungals, topical therapy and clinical debridement or nail avulsion. The newer oral anti-fungal agents like terbinafine hydrochloride, itraconazole, etc., have shown promising efficacies (~60%) in the treatment of fungal nail infections but has high relapse rates ranging from 25%-50%. In addition, adverse effects such as headache, risk of hepatotoxicity, potential drug-drug interactions limit the use of oral medications, leading to only 35%-65% of physicians prescribing oral therapy for the treatment of onychomycosis. Topical therapies range from the use of topical anti-fungal agents like terbinafine hydrochloride, itraconazole, etc., have shown promising efficacies (~60%) in the treatment of fungal nail infections but has high relapse rates ranging from 25%-50%.  

8.5 | Transungual iontophoretic drug delivery

The drug/device combination which uses iontophoresis to deliver high concentrations of medication directly to the site of action can greatly improve the treatment of onychomycosis. The literature demonstrating transungual iontophoresis dates back almost three decades. In 1986, James et al.  tried transungual delivery of prednisolone across thumbnail to achieve peak plasma levels of about one-third of that achieved by oral ingestion. Various studies were performed by Nair et al. on transungual delivery of terbinafine and in one of the studies, a 21- and 37-fold enhancement in terbinafine delivery was reported after 1 hour (0.5 mA/cm²) of iontophoresis in cadaverous human finger and toe nail respectively. A 150-fold increase in iontophoretic delivery of terbinafine through folded porcine epidermis as a model of proximal nail fold was demonstrated by Manda et al. An in vivo transungual iontophoretic experiment enhanced the flux of chloride and sodium ions by 8- and 27-fold, respectively, when compared to passive diffusion. Hao et al. demonstrated an enhancement of 10-fold in delivering ciclopirox olamine through cadaverous human nail plate iontophoretically, when compared to passive transport. Amichai et al. performed maiden clinical trials on iontophoretic delivery of terbinafine in human nails for the treatment of
onychomycosis and showed a significant decrease in frequency of patients showing presence of nail fungus at week 12 who wore 1% terbinafine iontophoretic patch (n=19) than compared to the passive treatment group wearing just 1% terbinafine patch (n=16). Amichai et al. attributed the enhanced efficacy of the iontophoretic patch to greater loading of drug in the nail plate, which helped in maintaining the levels of drug above the minimum inhibitory concentration (0.1-1 µg/cm²) required to be effective against fungal infection. Two of the patients (out of 19) reported local irritation and discomfort, whereas some of the patients reported tingling sensation. The iontophoretic patch treatment was considered to be overall tolerable and safe as none of the patients who reported tingling sensation or discomfort stopped the patch treatment.

9 | WHY IS THE NAIL FUNGAL INFECTION SO DIFFICULT TO TREAT?

Although the relative permeability per unit thickness of the nail plate is much higher than the stratum corneum, the thickness of the nail plate renders itself as a formidable barrier to anti-fungal drug delivery. In addition, the relapse rate of the patients treated for nail fungal infection can be as high as 50%. There might be multiple factors such as patients’ genetic predisposition, lifestyle and occupation which may lead to this high rate of fungal recurrence. Patients with compromised immune system or with concomitant diseases are more predisposed to onychomycosis and are unlikely to attain a lasting cure. The recurrence could also be associated either with failure to completely treat the infected nail or subsequent re-infection due to exposure to a different fungal strain. Nail fungal infection could also spread within household by being in close proximity with the infected family member. One of the most recently published studies demonstrated that members of the same household were infected with same strain of T. Rubrum. Even if the fungal infection is clinically eradicated, the patient still remains at risk of relapse due to exposure to fungal environment from public showers in gyms and hotel rooms. Thus, one must avoid being barefoot at places where the potential of fungal infection is high. As a preventative measure, one could spray the shoes with a topical anti-fungal spray or treat it with a commercially available UV shoe sanitiser after each use to avoid the re-infection.

10 | CONCLUSIONS

Although our knowledge of topical transungual delivery and nail permeability properties has enhanced considerably recently, much information is still unknown like the fine microstructure of the nail plate, structural mathematical model for predicting nail delivery and drug-keratin binding properties. The therapeutic approach to treat the nail fungal infections could possibly be achieved by a combination of topical therapy with physical or chemical enhancement methods. In addition, special precautionary steps must be taken to avoid the recurrence of the nail infection.

ACKNOWLEDGMENTS

This work was supported in part by a grant from the NIH (GM063559). The opinions expressed in this paper are those of authors and not endorsed by the NIH. The authors thank Mark Smith at Amway Corporation for providing assistance with nail SEM imaging.

DISCLOSURE STATEMENT

The author declares no conflict of interest.

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