# The effect of risk factors and etiology on the distribution of clinical cases with dermatomycoses

Kawther Mohammed Ali Hasan<sup>1</sup> and Majed Kadhum Aboud Al-Shibli<sup>2</sup>

<sup>1</sup>College of Science for Girls, Babylon University, Hilla, Iraq

<sup>2</sup>College of Education, Al-Qadisya University, Al-Qadisya, Iraq

Copyright © 2016 ISSR Journals. This is an open access article distributed under the *Creative Commons Attribution License*, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

**ABSTRACT:** The common cases of skin infections are dermatophytoses which included different clinical types. Aim of this study was evaluated the effect of risk factors such as age, gender, residency, chronic diseases and existence of domestic animals. In addition to identification of dermatophytic species by direct microscopic examination and cultured on Sabouraud's Dextrose Agar (SDA). Shows through the results that have been reached for clinical samples to 91 sample (46.66%) out of 195 samples were positive by direct microscopic examination, while the number of positive samples in culture 132 sample (67.69%). Tinea corporis was the most common (43.07%) of dermatophytoses, formed all kinds of Tinea exception of Tinea capitis ratio infected in females higher than in males. Also, all kinds of Tinea given a larger proportion of the infection of patients who live in rural areas from who live in urban areas. The highest rate of Tinea were in the age group 31-40 years. That most of the pathological cases of dermatophytoses associated with lack of chronic diseases of the 195 cases only 26 (13.33%) were infected with chronic diseases. The dominant species of fungus isolated was *E. floccosum* (34.84%).

**KEYWORDS:** Clinical samples, Dermatophytoses, Tinea, Risk factors, Direct microscopic examination.

## INTRODUCTION

Dermatomycoses were the common human diseases in which increased significantly in recent years, however that these diseases are not life threatening, but they are among the most prevalent diseases in the world and that cost millions of dollars to treat, According to Gräser *et al.*, (2008) that more than 500 million dollars annually is spent worldwide for drugs targeted against dermatophytoses, while some sources pointed out that in the United States alone, about more than 400 million dollars spent annually to treat dermatophytoses (Gaedigk *et al.* 2003; Jackson *et al.*,2006; White & Henn, 2007; Achterman and White, 2012). And therefore cause economic and public health problems together (Mirzahoseini *et al.*, 2009).

Although dermatophytoses is a nonfatal condition and not life threatening, this may have significant clinical consequences such as secondary bacterial infection, chronicity, disfigurement and therapeutic difficulties often necessitating long-term treatment, in addition to serving as a reservoir of infection especially onecomycosis (Malik *et al.*, 2009; Yamaguchi *et al.*,2009).

Although the penetration of tissue by the dermatophytes is usually quite superficial, adsorption of products from the fungi such as proteases leads to sensitization of the host, an event that is manifested by specific delayed hypersensitivity and other sorts of allergic responses especially zoophilic and geophilic dermatophytes are responsible for quite severe inflammatory reactions (Howard *et al.*, 2003; Zuzarte *et al.*, 2011).

The diseases caused by anthropophilic dermatophytes are commonly mild, while those caused by zoophilic dermatophytes are mostly sever and more inflammatory (Refai *et al.*,2013). This diseases can appear in different anatomical

areas of the body, so are classified according to the body areas that are affected such as Tinea capitis, Tinea coporis, Tinea cruris, Tinea pedis, Tinea unguium, Tinea faciei and others (Nejad *et al.*, 2007).

The aim of our study was to evaluate the risk factors and etiology effected on the dermatophytoses in clinical cases based on assess the prevalence of age and gender, the site of lesion, residency, existence of domestic animals and presence of chronic disease.

### MATERIAL AND METHODS

#### **SAMPLES COLLECTION**

A total of 195 specimens were collected from patients with dermatophytoses which including 60/195 (30.76%) specimens from male and 135/195 (69.23) specimens from female who clinically diagnosed by dermatologist. There are three types of specimens as follow: 20(10.25%) nails clips, 23(11.79%) hair fragment and 152(77.94%) skin scraps. The characteristics of the patient such as age, gender, the site of lesion, residency, existence of domestic animals and presence of chronic disease were collected in advance forms.

### DIRECT EXAMINATION AND CULTURE TEST

For direct microscopic examination, specimens were mounted in 10% potassium hydroxide (KOH), for identification of dermatophytes were cultured on Sabouraud's Dextrose agar (SDA) with chloramphenicol and cycloheximide and cultured at 26°C for up to 4 weeks. The identification of fungal agents were based on macro- and micromorphological characteristics. In addition, fungal identification was confirmed by the in vitro hair perforation test, urease production in Christensen's medium and vitamin requirements in Trichophyton agar media (Kannan *et al*, 2006).

Statistical analysis was performed by using the chi-square test to find the significant correlation at (p< 0.05) level .

### **RESULTS AND DISCUSSION**

The results of routine examinations revealed that 78/195 (40%) specimens were positive in both direct examination (KOH) and culture, 54/195 (27.69%) specimens were positive in culture and negative in KOH, and 13/195 (6.66) specimens were negative in culture and positive in KOH, while the specimens of negative in both direct examination and culture were 50/195 (25.64) (Figure 1).

Our results of routine examinations were consistent with the results of other workers (Kannen *et al.*, 2006; Madhavi *et al.*, 2011), the negative result of direct examination or culture may be due to little of specimens volume or the patients previously treated with antibiotics or take specimen from center of lesion which acquired local immunity but must be taking from margin of lesion and others reasons (Miline,1996; Cortez *et al.*, 2012), also the low positivity in direct examination of Tinea unguium is because of nail specimen take long time to dissolve and fungal element may not release and high positivity in culture could be due to use of selective media which does not allowed contaminates to grow (Mathur *et al.*, 2012).



Figure (1): Infection percentage of the direct examination and culture results

In this study Tinea corporis is the most common clinical form of dermatophytoses (43.07%) followed by T. cruris (18.46%), T. capitis (11.76%), T. unguium (10.25%), T. faciei (8.71%), T. manum (4.61%) and T. pedis (3.07%) respectively (Figure 2). These results were consistent with the other studies (Ellabib & Khalifa , 2001; Mathur *et al.*, 2012; Prasad *et al*, 2013) which showed that tinea corporis often have occupied the first place of dermatophytoses.



Figure (2): Percentage of clinical cases of dermatophytoses

According to the patients age it was found that statistically significant effect (P< 0.05) on the distribution of dermatophytoses, The results exhibited that the age 31-40 years had the highest frequency 50/195 (25.6%), while age group  $\geq$  60 years recorded the lowest frequency 2/195 (1.02%), the results was agreement with Pique *et al.* (2002) who found the highest frequency of dermatophytoses in age 30-40 years while the lowest frequency in age more than 70 years. The higher incidence in young males could be due to greater physical activity and increased sweating (Peerapur *et al.*, 2004). And the reason for less infection in higher ages can be justified as cellular immunity system perfection and the skin fatty acid augmentation (Nejad *et al.*, 2007).

Clinical type	Age (year)							Total
	≤ 10	11-20	21-30	31-40	41-50	51-60	≥ 60	No.(%)
	(%)	(%)	(%)	(%)	(%)	(%)	(%)	
Tinea corporis	8	16	25	24	5	5	1	84
	(4.1)	(8.2)	(12.8)	(12.3)	(2.56)	(2.56)	(0.51)	(43.07)
Tinea cruris	3	6	8	10	7	2	_	36
	(1.53)	(3.07)	(4.1)	(5.12)	(3.58)	(1.02)		(18.46)
Tinea capitis	19	2	_	1	1	_	_	23
	(9.74)	(1.02)		(0.51)	(0.51)			(11.79)
Tinea unguium	_	_	8	8	1	3	_	20
			(4.1)	(4.1)	(0.51)	(1.53)		(10.25)
Tinea faciei	4	4	2	3	2	2	_	17
	(2.04)	(2.04)	(1.02)	(1.53)	(1.02)	(1.02)		(8.71)
Tinea manuum	1	2	1	3	2	_	_	9
	(0.51)	(1.02)	(0.51)	(1.53)	(1.02)			(4.61)
Tinea pedis	_	1	1	1	1	1	1	6
		(0.51)	(0.51)	(0.51)	(0.51)	(0.51)	(0.51)	(3.07)
Total No.	35	31	45	50	19	13	2	195
	(17.9)	(15.9)	(23.1)	(25.6)	(9.74)	(6.66)	(1.02)	(100)

### Table (1): Distribution of dermatophytosis patients according to age

The effect of risk factors on clinical types of dermatophytosis was found that statistically significant effect (P < 0.05) (Table 2), the gender relationship were that all types of Tinea exception Tinea capitis were female infection highest frequency from male infection while infection rate in Tinea capitis (9.23%) male and (2.56%) female. Patients with tinea corporis recorded the highest frequency (13.33%) males and (29.74%) females while the lowest frequency in male was of Tinea unguium (0.51%) and in female was of Tinea pedis (2.05%), usually males infected with Tinea capitis more than females because of hormones that are play a role in increase, or may br the reason due to attributed to the easy implantation of spores because of short hair and frequency of sharing comb , brushes , and cups (Woldeamanuel *et al.*, 2005; Ilkit and Demirhindi , 2008; Younes *et al.*; 2012).

The influence of residence on distribution of dermatophytosis showed that 57.94% patients was from rural area while 42.05% patients was from urban area, in addition, all types of tinea were highest frequency in rural area exception tinea cruris was more prevalent in urban area (11.79%) compared with rural area (6.66%), the reason for elevation rural area infection may be its poor socioeconomic environmental that due to increase of infectious diseases especially among children (Inanir *et al.*, 2002).

Often of cases were not suffering from any chronic disease (86.6%) compared with cases that suffering from chronic disease (13.3%). Tinea corporis recorded the highest frequency including 6.15% cases with chronic disease followed by Tinea cruris while all of Tinea faciei and tinea manuum without chronic disease. This results refer to the disease with dermatophytes infected both immunocompetent or immunocompromised patients. Almost affect 10-20% of the world's population and it is possible to infect immunocompetent hosts, in many cases, the treatment methods are very difficult, or may re-infection after recovery (Recurrent) (Grumbt *et al.* 2011). In patients with diabetes, hyperglycaemia itself has been shown to decrease phagocytic activity further, thus dermatophytes infections (Bristow & Mak, 2009).

The patients in contact with domestic animals were 73.8% of cases while that no contact with domestic animals were 26.2%, Tinea corporis recorded the highest frequency including 31.8% cases with existence of domestic animals and 11.2% cases with absent of domestic animals. These results were consistent with Ayesh (2013) when he studied dermatophytosis in Gaza, Palestine. Increase of dermatophytes infections in Tinea unguium may be because majority of people are farmer by occupation and there are more chances of transmission of geophilic and zoophilic species (Mathur *et al.*, 2012).

Clinical	Gender (%)		Residency (%)		Chronic		Domestic		Total
type					disease (%)		animals(%)		No. (%)
	Male	Female	Rural	Urban	+	-	+	-	
Tinea	26	58	44	40	12	72	62	22	84
corporis	(13.33)	(29.74)	(22.56)	(20.51)	(6.15)	(36.9)	(31.8)	(11.2)	(43.07)
Tinea cruris	5	31	13	23	5	31	20	16	36
	(2.56)	(15.89)	(6.66)	(11.79)	(2.56)	(15.9)	(10.2)	(8.2)	(18.46)
Tinea	18	(2.56) 5	13	10	3	20	20	3	23
capitis	(9.23)		(6.66)	(5.12)	(1.53)	(10.2)	(10.2)	(1.53)	(11.79)
Tinea	1	(9.74) 19	13	(3.58) 7	4	16	15	5	20
unguium	(0.51)		(6.66)		(2.04)	(8.20)	(7.69)	(2.56)	(10.25)
Tinea faciei	5	(6.15) 12	17	_	_	17	15	2	17
	(2.56)		(8.71)			(8.71)	(7.69)	(1.02)	(8.71)
Tinea	3	(3.07) 6	(4.61) 9	_	_	9	9	_	9
manuum	(1.53)					(4.61)	(4.61)		(4.61)
Tinea pedis	2	(2.05) 4	(2.05) 4	(1.02) 2	2	4	3	3	6
	(1.02)				(1.02)	(2.04)	(1.53)	(1.53)	(3.07)
Total No.	60	135	113	82	26	169	144	51	195
	(30.76)	(69.23)	(57.94)	(42.05)	(13.3)	(86.6)	(73.8)	(26.2)	(100)

Table (2): Relationship	between clinical	type of dermatop	phytosis with r	isk factors
-------------------------	------------------	------------------	-----------------	-------------

Twelve of dermatophytes species were identified in specimens of examined patients, *E. floccosum* 46(34.8%) showed the highest frequency followed by *T. interdigetale* 19(14.4%) and *T. mentagrophytes* 18(13.6%) respectively (Table 3). *E. floccosum* was found to be predominant species in all types of Tinea cases exception Tinea capitis. These results were consistent with the other studies (Jha *et al.*, 2012; Abastabar *et al.*, 2013; Prasad *et al.*, 2013).

Table (3): Distribution o	f dermatophytosis	according to fungal isolates.
---------------------------	-------------------	-------------------------------

Fungal isolates	Tinea	Tinea	Tinea	Tinea	Tinea	Tinea	Tinea	Total No.
	corpoi	cruris	capitis	unguium	faciei	manum	pedis	(%)
E. floccosum	21	18	_	3	1	2	1	46 (34.8)
M. audouinii	2	-	_	_	-	_	-	2 (1.51)
M. canis	8	I	2		2	-		12 (9.09)
T. equinium	2	1	_	I	I		I	3 (2.27)
T. interdigetale	10	3	3	_	1	2	-	19 (14.4)
T. mentagrophytes	8	5	_	_	4	1	_	18 (13.6)
T. rubrum	2	1	2	_	-	_	2	7 (5.3)
T. schoenleinii	1	1	_	_	-	_	_	2 (1.51)
T. soudanense	2	-	_	_	-	_	_	2 (1.51)
T. tonsurans	_	-	2	_	-	_	_	2 (1.51)
T. verrucosum	7	-	3	_	2	_	_	12 (9.09)
T. violaceum	4	1	2	_	_	_	_	7 (5.3)
Total No.	67	30	14	3	10	5	3	132 (100)

## REFERENCES

- [1] Abastabar, M.; Rezaei-Matehkolaei, A.; Shidfar, M.R. & *et al.*, (2013). A molecular epidemiological survey of clinically important dermatophytes in Iran based on specific RFLP profiles of betatubulingene. Iranian J Publ Health, 42(9):1049-1057.
- [2] Achterman, R.R. and White, T.C. (2012). Dermatophyte virulence factors: identifying and analyzing genes that may contribute to chronic or acute skin infections. International J. Microbio., V(2012):1-8.
- [3] Ayesh, E.K. (2013). Improving the Diagnosis of Dermatophytes in Gaza Strip by using Nested PCR. A thesis of M.S., Eslamic Uni. of Gaza.

- [4] Bristow, I & Mak, M. (2009). Fungal foot infection: the hidden enemy? Wound UK 5(4):72-78.
- [5] **Cortez, A.A.; de Souza, J.V.B.; Sadahiro, A. & de Oliveira, J.A.A. (2012).** Frequency and aetiology of dermatophytosis in children age 12 and under in the state of Amazonas, Brazil . Revista Iberoamericana de Micologia 29(4): 223-226.
- [6] Ellabib, M.S.; Agaj, M.; Khalifa, Z. & Kavanagh, K. (2002). *Trichophyton violaceum* is the dominant cause of tinea capitis in children in Tripoli, Libya: results of a two year survey. Mycopathologia 153:145–147.
- [7] Gaedigk, A.; Gaedigk, R. & Abdel-Rahman, S.M.(2003). Genetic Heterogeneity in the rRNA Gene Locus of *Trichophyton tonsurans*. J. Clin. Microbio., 41(12): 5478–5487.
- [8] Gräser, Y.; Scott, J. & Summerbell, R.(2008). The new species concept in dermatophytes a Polyphasic approach. Mycopathologia V(2008):1-18.
- [9] Grumbt, M.; Monod, M. & Staib, P. (2011). Genetic advances in dermatophytes. FEMS Microbiology Letters 320:79-86.
- [10] Howard, D.H.; Weitzman, I. & Padhye, A.A. (2003). Onygenales: Arthrodermataceae. In:Pathogenic fungi in humans and animals.Marcel Dekker, Inc. New York, secod edition, 790 pp.
- [11] Ilkit, M. & Demirhindi, H. (2008). Asymptomatic dermatophyte scalp carriage: laboratory diagnosis, epidemiology and management. Mycopathologia, 165:61-71.
- [12] Inanir, I.; Sahin, M.T.; Gunduz, K.; Dinç, G.; Turel, A. & Ozturkcan, S. (2002). Prevalence of skin conditions in primary school children in Turkey: differences based on socioeconomic factors. Pediatric Dermatology, 19(9): 307-311.
- [13] Jackson, C.J.; Mochizuki, T. & Barton, R.C.(2006). PCR fingerprinting of *Trichophyton mentagrophytes* var. *interdigitale* using polymorphic subrepeat loci in the rDNA nontranscribed spacer. Journal of Medical Microbiology, 55: 1349–1355.
- [14] Jha, B.K.; Murthy, S.M. & Devi, N.L. (2012). Molecular identification of dermatophytosis by polymerase chain reaction (PCR) and detection of source of infection by restricted fragment length polymorphism (RFLP). J. College of Medical Sciences- Nepal, 8(4):7-15.
- [15] Kannan, P.; Janaki, C. & Selvi, G.S. (2006). Prevalence of dermatophytes and other fungal agents isolated from clinical samples. Indian Journal of Medical Microbiology, 24 (3):212-215.
- [16] Madhavi, S.; Rama, M.V. & Jyothsna, K. (2011). Mycological study of Dermatophytosis in rural population. Annals of Biological Research, 2(3):88-93
- [17] Malik, N.A.; Raza, N.& Nasiruddin (2009). Non-dermatophyte moulds and yeasts as causative agents in onychomycosis. Journal of Pakistan Association of Dermatologists, 19: 74-78.
- [18] Mathur, M.; Kedia, S.K. & Ghimire, R.B.K. (2012). Epizoonosis of Dermatophytosis: A Clinico Mycological Study of Dermatophytic Infections in Central Nepal. Kathmandu Univ Med J 2012;37(1):30-3.
- [19] Milne, L.J.R. (1996). Fungi In: Practical Medical Microbiology. 14th edn, New York: Churchill Livingstone. pp 695–720.
- [20] Mirzahoseini, H.; Omidinia, E.; Shams-Ghahfarokhi, M.; Sadeghi, G. & Razzaghi-Abyaneh, M. (2009). Application of PCR-RFLP to Rapid Identification of the Main Pathogenic Dermatophytes from Clinical Specimens. Iranian J. Publ Health, 38(1):18-24.
- [21] Nejad, S.B.; Khodaeiani, E. & Amirnia, M. (2007). A study of dermatophytosis infections in dermatology clinic of Sina hospital – Tabriz. Ege Tip Dergisi 46(1): 21 – 25.
- [22] Peerapur, B.V.; Inamdar, A.C.; Pushpa, P.V. & Srikant, B. (2004). Clinicomycological study of dermatophytosis in Bijapur. Indian Journal of Medical Microbiology, 22 (4):273-274.
- [23] **Prasad, N.; Mahapatra, A. & Chayani, N. (2013).** Changing trends in the fungal isolates from clinical specimens of suspected superficial mycosis. Indian Medical Gazette, 60-62.
- [24] Refai, M.; Heidy, A.E.; and Mahmoud , E. (2013). Monograph on dermatophytes. Department of Microbiology, Faculty of Veterinary Medicine, Cairo University. 75 pp.
- [25] White, T. & Henn, M. (2007). Genomic Determinants of Infection Competence in Dermatophyte Fungi. Dermatophyte Genome Steering Committee. 12 pp.
- [26] Woldeamanuel, Y.; Leekassa, R.; Chryssanthou, E.; Menghistu, Y. & Petrini, B. (2005). Prevalence of tinea capitis in Ethiopian school children. Mycoses, 48: 137-41.
- [27] Yamaguchi, M.U.; Silva, A.P.; Ueda-Nakamura, T.; Filho, B.P.; Silva, C.C. & Nakamura, C.V. (2009). Effects of a Thiosemicarbazide Camphene Derivative on *Trichophyton mentagrophytes*. Molecules, 14: 1796-1807.
- [28] Younes, A.K.H.; Mohamed, E.E.M.; Tawfik , K.M. and Ezzat, A.A. (2012). Tinia capitis in Assiut (Egypt). AAMJ 10(1): 45-54.
- [29] Zuzarte, M.; Gonçalves, M.J.; Canhoto, J. and Salgueiro, L. (2011). Antidermatophytic activity of essential oils. FORMA TEX : 1167- 1178.