Original Research Article

Treatment of dermatophyte toenail onychomycosis with itraconazole

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Received: 15 November 2016
Accepted: 09 December 2016

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ABSTRACT

Background: Onychomycosis is fungal infection of nail. The prevalence of onychomycosis varies across the world. It is lower in tropical countries (3.8%) than in sub-tropical countries and countries in the temperate zone (23%). The risk of onychomycosis is higher in persons with diabetes mellitus and human immunodeficiency virus infection. Systemic antifungals are recommended for treatment. Topical antifungals are less effective. Different cure rates have been reported by different authors for terbinafine, itraconazole and fluconazole therapy. A variety of regimens (continuous or pulse) have also been suggested.

Methods: Here in this study we report a prospective, observational, cohort study of pulse itraconazole therapy in toenail onychomycosis caused by dermatophytes over a period of one year in patients with SCIQ scores between 6 to 9.

Results: It was observed that mean age of patients was 38.96 years. 57.27 % study subjects had complete clinical improvement to oral itraconazole three pulses regimen. One pulse consisted of 200 mg twice per day given for one week, followed by a three weeks interval. Thus, this regimen was repeated thrice.

Conclusions: This study shows the substantial benefit with itraconazole pulse regimen in toenail onychomycosis. The outcome this study is comparable with previously published data and may encourage the usage of itraconazole pulse regimen for the treatment of dermatophyte toenail onychomycosis in adults.

Keywords: Itraconazole, Onychomycosis, Pulse therapy

INTRODUCTION

Onychomycosis is the most common fungal infection of the nail bed, matrix and/or plate, representing up to 50% of all nail disorders. It causes discolouration, disfigurement and separation of nail from nail bed. Onychomycosis affects toenails more than fingernails because of their slower growth, reduced blood supply and frequent confinement in the dark, moist environment. Onychomycosis can be caused by dermatophytes, nondermatophyte molds (NDM) and candida species. Dermatophytes of genera Trichophoton and Epidermophyton are capable of invading nails. These breakdown and utilize keratin as a source of nitrogen. Dermatophytes responsible for onychomycosis are Trichophoton rubrum, Trichophoton mentagrophytes, Trichophoton tonsurans, Trichophoton violaceum, Trichophoton schoenleinii, Trichophoton veruucusum and Epidermophyton floccosum. Nondermatophyte molds (fungi) include Aspergillus spp., Fusarium spp., Penicillium spp., Acremonium spp., Alternaria spp., Cladosporium carrion and Geotrichum candidum.¹²

Accurate diagnosis of onychomycosis involves clinical and microscopic examination (for identification of fungal hyphae, pseudohyphae or spores) and culture (identification of organism). Histological evaluation using Periodic- Schiff staining increases the sensitivity for detecting the infection (identify organism). Treatment is aimed at the eradication of the causative
organism and return to normal appearance of the nail. Systemic antifungals are the most effective treatment. Topical therapy with ciclopiroprin nail lacquer, amorolfine nail lacquer, and lacquer containing encecalin extract of Ageratina pichichense, efinaconazole solution or tavalaborol solution is less effective. To enhance the effectiveness of topical drugs attempts have been made to add penetration enhancer or incorporate the drug into transferome lipid vesicle. Surgical or chemical removal of infected nail parts is useful. It facilitates the penetration of antifungal drugs and eradicates reinfection sites so as to prevent recurrence and relapse. Laser therapy, photodynamic therapy, iontophoresis, ultrasound therapy, UVC radiation and psoralen with UVA (PUVA) show promise for onychomycosis treatment.1,4

Azoles group of oral antifungal drugs are used for the treatment of dermatophyte onychomycosis. Azoles are broadly classified into two groups - imidazoles and triazoles. They share the same antifungal spectrum and mechanism of action. The triazoles are metabolized more slowly and also have lesser effect on human sterol synthesis than imidazoles. The itraconazole, fluconazole, terconazole, voriconazole, posaconazole and isavuconazole are triazoles. The Imidazoles include clotrimazole, miconazole, ketoconazole, econazole, butaconazole, oxiconazole, sertaconazole and sulconazole. Terbinafine a synthetic allylamine is also effective in nail onychomycosis.3,6

Azoles inhibit the growth of fungi by inhibiting a microsomal cytochrome P450 (CYP) enzyme 14-alpha-sterol demethylase. Inhibition of this enzyme impairs the biosynthesis of ergosterol for cytoplasmic membrane and leads to accumulation of 14-alpha-methyl-sterols which disrupt close packing of acyl chains of phospholipids thus impair the function of membrane-bound enzyme systems (ATPase and electron transport system enzymes). Azoles may also directly increase permeability of fungal cytoplasmic membrane.

Itraconazole is a synthetic triazole. It is an equimolar racemic mixture of four diastereo - isomers, each possessing three chiral centers. It is available in a capsule form. It is best absorbed in fed state. The half-life of itraconazole at a steady state is approximately 30-40 hours. It is metabolized in the liver. It is usually prescribed for toenail onychomycosis as 200 mg once daily orally for three months - continuous therapy, or 200 mg twice daily orally for one week out of each month for three months- so called three pulse therapies. Retention of active drug in nail keratin permits for 28 weeks the intermittent treatment. Adverse drug reactions include nausea, vomiting, diarrhoea, abdominal discomfort, increased serum aminotranferase, increased serum triglycerides, hypokalemia, rash, hypertension, prolonged Q-T interval, ventricular dysfunction, congestive heart failure, lower limb edema, adrenal deficiency, rhabdomyolysis and stevens-johnson syndrome (rare).

The clinical types of onychomycosis of toenail are distal lateral subungal onychomycosis (DLSO), proximal subungal onychomycosis (PSO), total dystrophic onychomycosis (TDO) and Enconlyx sublingual onychomycosus (ESO).

The aim of present study is to evaluate the effectiveness of itraconazole, three pulse therapies, for the treatment toenail onychomycosis caused by dermatophytes, in adults.

METHODS

The present prospective, observational study was conducted on patients who attended Out Patient Department (OPD) of the Department of Skin and VD, Darbhanga Medical College and Hospital, Darbhanga during period January 2015 to December 2015. One hundred seventy nine patients with clinical signs of persistent onychomycosis of toes were enrolled on for the study. Primary criteria for the clinical diagnosis of onychomycosis included white/yellow or orange/brown streaks or patches in or beneath the nail, and secondary criteria included lateral onycholysis, subungal hyperkeratosis and nail plate thickening. One hundred seventeen adults were included in the study group. The evaluation of disease severity was done on the basis of SCIO onycho-index and calculated by electronic calculator in the link http:www.onychoindex.com. Written consent was obtained from each patient. Drugs were branded and purchased by patients. The study was approved from institutional ethics committee.

Inclusion criteria

- Adults patients (age >18 years)
- Males and females (non-pregnant and non-lactating females)
- Clinical signs of onychomycosis of at least one great toe nail
- Positive microbiological evidence of fungal elements in nail specimen
- Positive culture for dermatophyte in nail specimen

Exclusion criteria

- Females who are pregnant, lactating or likely to be pregnant
- Negative potassium hydroxide stain
- Negative mycological culture for dermatophytes consistent with onychomycosis
- Onychomycosis not caused by dermatophytes
- Immuno compromised either because of concomitant diseases e.g. HIV infection, or ongoing treatment e.g. chemotherapy
- Patients suffering from psoriasis, diabetes mellitus, renal failure, acute porphyria, ventricular dysfunction, congestive heart failure, abnormal liver function tests and haematological count abnormalities
Patients with history of serious liver disease, congestive heart failure, intake of oral antifungal drugs within 6 months prior to study, hypersensitivity to antifungal drugs

Patients on concomitant therapy with drugs requiring P450 enzyme system.

Procedures

Nail specimen of each patient was collected from the toenail most affected by using an electric dental drill, after cleaning the area with 70% isopropyl alcohol to prevent contamination. The specimen was placed on a slide and a solution 20% potassium hydroxide and 40% dimethyl sulfoxide was dropped on the specimen. The prepared slide was heated for two to five minutes by an electric hot plate set at sixty to eighty degree Celsius. The specimen was then examined by a microscope under x100 magnification for the presence of fungal elements (hyaline septate hyphae and/or arthroconid). The nail specimen of each patient was also sent to the laboratory in a sterile container for culture of dermatophytes. The specimen was inoculated into three sets of test tubes - first containing Sabouraud’s dextrose agar (SDA) with 0.05% chloramphenicol, second Sabouraud’s dextrose agar (SDA) with chloramphenicol and 0.5% cyclohexidine, and third Dermophyte test medium (DTM). Growth of dermatophytes in the media was observed regular once a week for 1 to 3 weeks. If no growth was detected after 3 weeks, specimen was considered as negative for dermatophytes. Dermatophyte species identification was done on the basis of characteristics colony morphology, growth rate (fast to very slow), characteristics microscopic features of various species of dermatophytes visible in the culture, supplemented by urease test, hair perforation test, slide culture and rice grain test. Medical data collected for each patient included age, gender, duration of onychomycosis, clinical types and SCIO score (scoring clinical index for onychomycosis score). SCIO score was calculated on the basis of three clinical factors

- Clinical form of onychomycosis (DLSO/ PSO/ SWO/ TDO) which indicated location of fungus in nail
- Depth of nail involvement (onychomycosis localized to first one third / two third / more than two third nail plate length)
- Thickness of subungual hyperkeratosis (absent or nor more than one mm/ one to two mm/ more than two mm).

SCIO score range from 1 to 30 and higher score indicates more severity. SCIO score 1 corresponds to SWO or marginal DLSO, while SCIO score 30 corresponds to whole nail DLSO with hyperkeratosis more than 2 mm.

Treatment regimen

Patients received a pulse regimen of itraconazole (200 mg twice daily) given after meals during the first week of each month, for three months. Patients were requested to visit OPD of the Department of Skin and VD, DMCH once a month for clinical check-up. At each visit clinical examination of nail and different systems of body was done.

Complete blood count, liver function tests (serum bilirubin, serum aspartate aminotransferase, serum alanine aminotransferase, serum alkaline phosphatase, and serum 5’ nucleotidase and serum gamma-glutamyl transpeptidase), serum triglyceride, serum potassium, echocardiography and electrocardiography were carried out at the baseline and two weeks after the each three pulses. Mycological microscopy and cultures were performed at baseline and twelve months after the start of treatment. Effectiveness (clinical cure and mycological cure) was evaluated twelve months after the start of treatment.

Primary outcome measure

Clinical cure

Defined as 100% absence of clinical signs of onychomycosis, evident by normal appearing nail.

Secondary outcome measure

Mycological cure

Defined as absence of dermatophyte on microscopy and culture in nail specimen.

Patients were asked to report any adverse drug reaction (ADR) if they occur. Assessment of reported ADR for causality, severity and preventability were undertaken using Naranjo’s causality, Hartwig and Siegel and Schumock and Thornton scales respectively.8-10

Statistical analysis

Statistical analysis was performed using SPSS version 20 for windows, (SPSS, Inc, Chicago, Illinois, United States of America). Data was expressed in Mean±standard deviation (SD).

RESULTS

One hundred seventeen adult patients suffering from dermatophyte onychomycosis of toenail were included in the study group. Seventy one were females and forty six males. Seventy four patients had distal lateral subungual onychomycosis (DLSO), thirty nine total dystrophic onychomycosis (TDO), two superficial white onychomycosis (SWO) and two proximal subungual onychomycosis (PSO). Trichophoton rubrum was isolated.
in seventy four cases, Trichophoton mentagrophytes in thirty one, Trichophoton tonsurans in six, Trichophoton violeceum in three and Trichophoton scholelini in three cases. Mean age of study group patients was 38.96 years; mean duration of disease was 6.31 years (Table 1).

Clinical cure was noted in sixty seven patients. Mycological cure was noted in seventy three patients. Three patients experienced nausea, two complained of vomiting and one experienced rash during first pulse therapy.

Table 1: Baseline characteristics of study subjects.

| Characteristics                  | n (%)     |
|----------------------------------|-----------|
| Gender                           |           |
| Male                             | 46 (39.32)|
| Female                           | 71 (60.68)|
| Onychomycosis types              |           |
| Distal lateral subungal onychomycosis (DLSO) | 74 (63.25) |
| Total dystrophic onychomycosis (TDO) | 39 (33.33) |
| Superficial white onychomycosis (PWO) | 2 (1.71)  |
| Proximal subungal onychomycosis (PSO) | 2 (1.71)  |
| Dermatophytes                    |           |
| Trichophoton rubrum              | 74 (63.25)|
| Trichophoton mentagrophytes      | 31 (26.50)|
| Trichophoton tonsurans           | 6 (5.13)  |
| Trichophoton violeceum           | 3 (2.56)  |
| Trichophoton scholelini          | 3 (2.56)  |
| Characteristics                  | Mean (SD) |
| Age (year)                       | 38.96 (12.13)|
| Duration of disease (year)       | 6.31 (2.29)|

n = number of patients, % = percentage, SD = standard deviation.

Table 2: Outcome variables.

| Outcome                          | n (%)     |
|----------------------------------|-----------|
| Primary                          |           |
| Clinical cure achieved           | 67 (57.26)|
| Clinical cure not achieved       | 23 (19.66)|
| Secondary                        |           |
| Mycological cure achieved        | 73 (62.39)|
| Mycological cure not achieved    | 17 (14.53)|

n = number of patients, % = percentage.

Table 3: Evaluation of disease severity.

| Duration                          | SCIO score mean (SD) |
|-----------------------------------|-----------------------|
| At baseline                       | 7.79 (0.89)           |
| 3 months after start of treatment | 3.56 (1.19)           |
| 12 months after start of treatment| 0.51 (1.23 )          |

SCIO = Scoring clinical index for onychomycosis score, SD = Standard deviation.

They received symptomatic treatment, were improved but discontinued treatment. Abnormal liver function tests or haematological reports were not observed in any patient during follow up.

Table 4: Treatment response on onychomycosis with itraconazole pulse therapy.

| Onychomycosis clinical types | At baseline n (%) | During 1 week n (%) | Follow up 12 months n (%) | Treatment response |
|------------------------------|-------------------|---------------------|---------------------------|--------------------|
| DLSO                         | 74 (63.25)        | 72 (61.54)          | 63 (53.85)                | Clinical cure 60 (51.28) |
| TDO                          | 39 (33.33)        | 36 (30.77)          | 24 (20.51)                | Mycological cure 10 (8.55) |
| SWO                          | 2 (1.71)          | 1 (0.85)            | 1 (0.85)                  |                    |
| PSO                          | 2 (1.71)          | 2 (1.71)            | 1 (0.85)                  |                    |
| Total                        | 117 (100)         | 111 (98.87)         | 90 (76.92)                | 67 (57.26)         |

DLSO = Distal lateral subungal onychomycosis; TDO = Total dystrophic onychomycosis; SWO = Superficial white onychomycosis, PSO= Proximal subungal onychomycosis; At 1 week, six patients discontinued due to adverse drug reactions; 12 months after start of treatment. 21 patients were lost during follow up.

Table 5: Adverse drug reactions (ADR) observed during study.

| ADR               | Casualty score (Naranjo’s scale) | Severity grade (Hart wig and Siegel scale) | Preventability grade (Schumock and Thornton scale) |
|-------------------|---------------------------------|------------------------------------------|--------------------------------------------------|
| Nausea            | 2 (Possible)                    | Moderate (Level 3)                       | Not preventable                                  |
| Nausea            | 3 (Possible)                    | Mild (Level 2)                           | Not preventable                                  |
| Nausea            | 3 (Possible)                    | Mild (Level 2)                           | Not preventable                                  |
| Vomiting          | 2 (Possible)                    | Moderate (Level 3)                       | Not preventable                                  |
| Vomiting          | 2 (Possible)                    | Mild (Level 1)                           | Not preventable                                  |
| Rash              | 3 (Possible)                    | Mild (Level 1)                           | Not preventable                                  |
After 3 months

After 12 months

In mycological itraconazole

DISCUSSION

Mean (SD) SCIO score decreased from 7.79 (0.89) at baseline to 0.51 (1.23) twelve months after the start of treatment (Table 2), (Table 3), (Figure 1), (Table 4), (Table 5).

Figure 1: Evaluation of disease severity.

The present study observed lower clinical and mycological cure rates with pulse therapy regimen with itraconazole, than reported by many previous studies except one. The present study observation is similar to other studies as regards to the safety. Only 5.13% patients discontinued the study due to ADR. The only limitation of the present study is that long-term follow up has not been done to investigate for clinical and mycological recurrence.

CONCLUSION

Itraconazole oral pulse therapy is effective and safe and can be recommended for the treatment of dermatophyte toenail onychomycosis in adults. Further trials are required to evaluate number of pulse therapies required for dermatophyte toenail onychomycosis of different SCIO scores.

ACKNOWLEDGEMENTS

Authors would like to thanks all the participants of the study.

Funding: No funding sources
Conflict of interest: None declared
Ethical approval: The study was approved by the Institutional Ethics Committee

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Cite this article as: Giri VP, Gupta SK, Giri OP, Kirti V. Treatment of dermatophyte toenail onychomycosis with itraconazole. Int J Basic Clin Pharmacol 2017;6:70-5.