



Seroepidemiology of *Strongyloides stercoralis* amongst immunocompromised patients in Southwest Iran



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ABSTRACT

Strongyloidiasis is a life-threatening parasitic infection, especially in immunosuppressed patients, with death often occurring within several days. The disease has a worldwide distribution and is endemic in tropical and subtropical regions. Therefore, this study was conducted to investigate seroepidemiology of *Strongyloides* infection amongst immunocompromised patients in Southwest Iran. This cross-sectional study was conducted amongst a population of immunocompromised patients who were referred to health care or hospital referral centres in Ahvaz, Southwest Iran. Serum samples were tested by an enzyme immunoassay for anti-IgG *Strongyloides* antibody. Anti-*Strongyloides stercoralis* antibody was detected in 39 of 270 immunocompromised cases, yielding a prevalence of 14.4%. No significant differences were indicated in terms of gender, age, or type of immunocompromised disorder with anti-*Strongyloides stercoralis* antibody levels. In conclusion, our results demonstrated high seroepidemiology of infection with this parasite in the region. Therefore, it appears immunocompromised patients should be tested for this infection using sensitive tests. However, current research underscores that strongyloidiasis must not be neglected, and further assessments in high risk population are warranted.

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1. Introduction

Strongyloidiasis is a condition caused by *Strongyloides stercoralis*, which may be established in humans for extended periods of time because of its auto-infective life cycle. It may become life-threatening especially in immunocompromised hosts (Grove, 1996; Sharifdini et al., 2014). Strongyloidiasis symptoms may vary from subclinical in acute and chronic infection to severe and fatal in hyper infection syndrome and disseminated strongyloidiasis which, if untreated, may approach mortality rates of up to 90% (Fernando et al., 2014; Annette et al., 2009).

The disease has worldwide distribution and is endemic to tropical and subtropical regions, such as Latin America, Asia, parts of Africa and the southeastern United States (Genta, 1989; Steinmann et al., 2007; Scha et al., 2013; Román-sánchez et al., 2003). Strongyloidiasis has also been reported in most parts of Iran with a higher prevalence amongst immunocompromised populations (Zali et al., 2004; Nilforoushan et al., 2007; Meamar et al., 2007a; Froutan et al., 2008; Keiser & Nutman, 2004).

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Recent developments in medical care have resulted in increasing numbers of immunocompromised individuals living for longer periods, which increases their risk of exposure for a longer duration, (Roseman et al., 2013; Paul et al., 2010). Parasitological diagnosis is often time-consuming, as *Strongyloides* larvae in the stool are frequently absent or present in very small numbers (Afzal et al., 2001). Recently, serological assays have been demonstrated to have reliable sensitivity and specificity for antibody detection (Salvador et al., 2014; Bisoffi et al., 2014; Lindo et al., 1995). Thus, the current study was conducted to investigate the seroepidemiology of *Strongyloides* infection amongst immunocompromised patients in Southwest Iran.

2. Materials and methods

2.1. Study population and sample collection

This cross-sectional study was carried out amongst a population of immunocompromised patients referred to health care or hospital referral centres in Ahvaz, Southwest Iran. Blood samples were collected from participants from March–July 2015. Participants suffered from leukaemia, HIV infections and different types of cancer. Demographic information including gender, age and immunocompromised disorders were collected from hospital and medical official documents. All blood specimens were centrifuged, and the serum separated and frozen at $-20\text{ }^{\circ}\text{C}$ prior to analysis for parasite-specific IgG antibodies pending examination by an enzyme-linked immunosorbent assay (ELISA) for *S. stercoralis*.

2.2. Serology assay

An enzyme immunoassay for the diagnosis of human strongyloidiasis (Bordier affinity products SA, Switzerland) was used. The method was carried out according to the manufacturer's instructions. Briefly, wells pre-coated and sensitised with *Strongyloides* somatic larval antigens were blocked with PBS-Tween for 15 min. Wells were washed with PBS-Tween then diluted sera [1:200] were added and incubated at $37\text{ }^{\circ}\text{C}$ for 15'. Serum was removed and the plates were washed four times. Alkaline phosphatase was added, plates washed, followed by the substrate. Finally, the reaction was stopped using K_3PO_4 . Well absorbances were measured at 405 nm. The cutoff point was used to discriminate optimally between the sera of clinically documented strongyloidiasis cases and normal human sera. The ELISA had a sensitivity of 88% and a specificity of 94%. In addition to positive and negative controls from manufacturer using a local strongyloidiasis serum, readings were confirmed by stool examination, which was also included in each run of ELISA assay.

3. Results

The study population included 166 (61.5%) AIDS-defining illness cases, 36 (13.3%) leukaemia cases and 68 (25.2%) other cancer cases. *S. stercoralis* antibody was detected in 39 immunocompromised individuals, yielding a prevalence of 14.4%. The frequency of seropositivity and immunosuppressive characteristics of the study samples are shown in Table 1. As indicated, the highest rate of antibody detection (15.7%) was found in HIV⁺ cases.

No significant differences were recorded in terms of gender, age, or immunosuppressive disorder with *anti-S. stercoralis* antibody levels (Table 2).

Additionally, the gender distribution of seropositive females and males has been reported to be 16.7% and 13.3%, respectively (Table 3).

4. Discussion

Strongyloidiasis is a parasitic infection characterised by persistent infection, dissemination and the development of potentially fatal disease. This infection has a worldwide distribution, although no precise estimate is available. The present study shows the seroepidemiological prevalence of *S. stercoralis* infection amongst immunocompromised cases in Southwest Iran to be 14.4%. Meamar et al. conducted an investigation on strongyloidiasis in 781 HIV⁺/AIDS patients by the stool examination technique. According to their results, two were determined to be infected (Meamar et al., 2007b). One report showed a 4.9% *S. stercoralis* infection rate amongst rural area of Mazandaran province in Iran during 2007 (Kia et al., 2007). Additional research indicated,

Table 1
Frequency of seropositivity of *S. stercoralis* infection amongst immunocompromised patients, Southwest Iran.

| Immunocompromised cases | Number | IgG positive | |
|-------------------------|--------|--------------|-----------|
| | | No | Frequency |
| HIV | 166 | 26 | 15.7% |
| Leukaemia | 36 | 4 | 11.1% |
| Cancer | 68 | 9 | 13.2% |
| Total | 270 | 39 | 14.4% |

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