RESEARCH ARTICLE

Imported strongyloidiasis: Data from 1245 cases registered in the +REDIVI Spanish Collaborative Network (2009-2017)

Fernando Salvador†*, Begoña Treviño2, Sandra Chamorro-Tojeiro3, Adrián Sánchez-Montalvá1, Juan María Herrero-Martínez4, Azucena Rodríguez-Guardado5, Núria Serre-Delcor6, Diego Torrés6, Josune Goikoetxea7, Zuriñe Zubero8, María Velasco9, Elena Sulleiro1, Israel Molina1, Rogelio López-Vélez3, José Antonio Pérez-Molina1, on behalf the +REDIVI Collaborative Network1

1Vall d’Hebron University Hospital, Universitat Autònoma de Barcelona, PROSICS Barcelona, Barcelona, Spain, 2 Tropical Medicine and International Health Unit Drassanes-Vall d’Hebron, PROSICS Barcelona, Barcelona, Spain, 3 Ramón y Cajal University Hospital, Madrid, Spain, 4 Hospital Universitario 12 de Octubre, Madrid, Spain, 5 Hospital Universitario Central de Asturias, Oviedo, Spain, 6 Hospital General Universitario de Alicante, Alicante, Spain, 7 Hospital Universitario Cruces, Barakaldo, Spain, 8 Hospital de Basurto, Bilbao, Spain, 9 Hospital Universitario Fundación Alcorcón, Madrid, Spain

†Membership of the +REDIVI Collaborative Network is listed in the S1 Appendix.

* fmsalvad@vhebron.net

Abstract

Background
Imported strongyloidiasis is increasingly being diagnosed in non-endemic areas. The aim of this study was to describe the epidemiological, clinical and microbiological characteristics of patients with imported strongyloidiasis in Spain.

Methodology
This is an observational retrospective study that included all patients diagnosed of strongyloidiasis registered in the +REDIVI Spanish Collaborative Network (2009-2017). Demographic, epidemiological and clinical information was collected from the +REDIVI database, and extra information regarding microbiological techniques, treatment and follow-up was requested to participant centers.

Findings
Overall, 1245 cases were included. Most of them were immigrants (66.9%), and South America was the most frequent area of origin. Detection of larvae in stool samples was observed in 21.9% of the patients, and serological tests allowed making the diagnosis in the rest of the cases. Eosinophilia was present in 82.2% of cases. Treatment with ivermectin (compared with albendazole) was the most strongly associated factor to achieve the cure (OR 2.34).
Conclusions
Given the long latency of the infection and the risk of developing a severe presentation, screening of S. stercoralis infection should be mandatory in patients coming from or had traveling to endemic areas, especially in those with immunosuppressant conditions.

Author summary
Strongyloidiasis, the parasitic disease caused by the nematode Strongyloides stercoralis, is being increasingly diagnosed in immigrant population in Spain. In the present study we describe the clinical, epidemiological and microbiological profile of 1245 cases of imported strongyloidiasis registered in the +REDIVI Spanish Collaborative Network. Patients were mostly immigrants, and the majority of them were coming from South America. Serological tests allowed the diagnosis in most of the cases, and detection of larvae in stool samples was associated with male gender and presence of eosinophilia. Ivermectin was associated with higher probability of cure compared with albendazole. Given the risk of developing a severe presentation, screening of S. stercoralis infection should be mandatory in patients coming from or had traveling to endemic areas, especially in those with immunosuppressant conditions.

Introduction
Strongyloidiasis is caused by the soil-transmitted nematode Strongyloides stercoralis [1]. The parasite is globally distributed in the tropics and occurs in some parts of the sub-tropics, although it has also been reported in more temperate climates, like southern Europe and North America [2]. It has been recently estimated that 370 million people are infected by the parasite; however, this prevalence is most probably underestimated, since the majority of the studies are focused on other soil-transmitted helminthes, and the microbiological techniques commonly used have low sensitivity for S. stercoralis [3, 4].

S. stercoralis infection is most often asymptomatic, but patients may present with gastrointestinal, cutaneous, and respiratory symptoms [2]. This parasite has the unique ability to persist in the human host for decades without the need of exogenous reinfection due to its autoinfection life cycle. Nevertheless, some immunosuppressant conditions, such as corticosteroid therapy, solid organ or bone marrow transplantation and Human T-lymphotropic virus 1 (HTLV-1) infection, may amplify this autoinfective cycle, leading to severe presentations such as the S. stercoralis hyperinfection syndrome and the disseminated strongyloidiasis [5, 6].

Due to increase of migrant flows and international travels during the last decades, imported strongyloidiasis is being increasingly diagnosed in Spain, as well as in other non-endemic areas, becoming a worldwide public health challenge [7–11]. The aim of the present study was to describe the epidemiological, clinical and microbiological characteristics of patients with imported strongyloidiasis registered in the +REDIVI Spanish Collaborative Network, as well as to identify potential factors related to parasite burden and treatment success.

Material and methods
Study design and study population
This is an observational retrospective study that included all patients diagnosed of strongyloidiasis registered in the +REDIVI Collaborative Network from January 2009 to February 2017.
This Spanish network includes 22 centers which share a common online database to register epidemiological and clinical information of immigrants and travelers who attend these centers. The database includes: demographic information (date of birth, gender, and country of birth), epidemiological data (country of origin for immigrants, travel characteristics), and clinical information (main reason for consultation, and diagnosis). +REDIVI includes 4 types of patients: immigrants (people living in Spain and born in other country), Visiting Friends and Relatives (VFR)-immigrants (immigrants who have travelled to their country of birth to visit friends and relatives), VFR-travelers (people born in Spain who have travelled to their first-degree relative's country of birth), and travelers (people born in Spain who return from international travels).

All participant centers were requested to contribute with extra information from the included patients: HTLV-1 co-infection, eosinophil cell count at diagnosis, microbiological techniques performed and their results, treatment information, and follow-up controls up to two visits.

**Microbiological techniques**

Stool microscopic examination was performed using the Ritchie’s formalin-ether technique in all centers. The specific fecal culture for *S. stercoralis* larvae was performed using the agar plate or charcoal technique. Detection of serum anti-*S. stercoralis* IgG was performed through different enzyme-linked immunosorbent assay (ELISA)-based commercial kits depending on the center: SciMedx *Strongyloides* serology microwell ELISA (SciMedx Corporation, Denville, NJ, United States), NovaLisa Strongyloides (NovaTec Immunodiagnostica GmbH, Dietzenbach, Germany), *Strongyloides* IgG IVD-ELISA (DRG Instruments GmbH, Marburg, Germany), AccuDiag *Strongyloides* IgG ELISA Kit (Diagnostic Automotion/Cortez Diagnostics Inc, CA, United States). All serological tests were performed and cut-off of the optical density values to determine a positive result was established following manufacturer instructions.

**Definitions**

Eosinophilia was defined as eosinophil cell count $\geq 0.45 \times 10^9$ cells/L and/or a percentage $\geq 5%$: mild ($<1.0 \times 10^9$cells/L), moderate ($1.0–3.0 \times 10^9$cells/L), and severe ($>3.0 \times 10^9$cells/L). The *S. stercoralis* infection diagnosis was classified into three groups: definite (detection of larvae by any parasitological technique), probable (positive serological result and presence of eosinophilia), and possible (positive serological result without presence of eosinophilia). We considered severe clinical presentation: hyperinfection syndrome (infection confined to lungs and gastrointestinal tract, but symptoms of severe disease related to elevated number of larvae) and disseminated strongyloidiasis (larvae found in any organ other than the lungs and gastrointestinal tract). In order to evaluate the treatment outcome, four categories were considered. The “cure” criteria was established when patients had negative conventional methods (microscopic stool examination and/or fecal culture) after treatment, disappearance of the eosinophilia, and negative serology (or at least a 60% decrease in the OD ratio) during follow-up (independently of the time they reached this situation). We established “probable cure” criteria when the cure criteria were reached but eosinophilia persisted (because it could be to other cause) or when in the absence of serologic control, patients had negative conventional methods after treatment and disappearance of the eosinophilia. “Failure” was considered if larvae were detected after the treatment or when the cure or probable cure criteria regarding serology were not reached during follow-up (with a minimum of 6 months of follow-up). The outcome “not enough information” was considered when the patients had negative conventional methods after
treatment, but cure or probable cure criteria regarding eosinophilia and serology were not reached during follow-up (with a follow-up period less than 6 months).

**Statistical analysis and ethical issues**

The STROBE statement guidelines were used to improve the quality of the study [12]. Procedures were performed in accordance with the ethical standards laid down in the Declaration of Helsinki as revised in 2013, and the study protocol was approved by the Ethical Review Board of the Vall d’Hebron University Hospital (Barcelona, Spain). The Ethical Review Board was consulted, and agreed that informed consent was not necessary given the retrospective nature of the study and the anonymization of the information.

Categorical data are presented as absolute numbers and proportions, and continuous variables are expressed as means and standard deviations (SD) or medians and interquartile ranges (IQR) depending on the distribution. The Kolmogorov-Smirnov test was used to evaluate the normal distribution of variables. The $\chi^2$ test or Fisher exact test, when appropriate, was used to compare the distribution of categorical variables, and the t-Student test for continuous variables. Multivariate logistic regression analysis was made to identify factors associated with confirmed diagnosis and cure. Variables were entered in the model if $p<0.20$ on univariate analysis. Results were considered statistically significant if the 2-tailed $P$ value was $<0.05$. SPSS software for Windows (Version 19.0; SPSS Inc, Chicago, IL, USA) was used for statistical analyses.

**Results**

Overall, 12796 patients were registered in +REDIVI from January 2009 to February 2017. Of these, 1279 patients had the *S. stercoralis* infection diagnosis. After revising these cases, 34 patients were withdrawn due to erroneous codification or duplication; hence, resulting in 1245 (9.7%) patients with *S. stercoralis* infection diagnosis (Fig 1). The mean age at the time of diagnosis was 38.3 (SD 12.9) years, and 590 (47.4%) were men (see Table 1). Most of them were immigrants (833 patients, 66.9%), and 80 (6.4%) patients had some kind of immunosuppressive condition. The main reasons for consultation were “laboratory test alteration” (41.1%) and "health screening" (39.4%). Only 5 (0.4%) patients had the diagnosis of hyperinfection syndrome: two patients with autoimmune diseases who were receiving immunosuppressive therapies (including corticosteroids), one patient with HTLV-1 infection, one patient with HIV infection, and one patient without any immunosuppressant condition. Six hundred and five (48.6%) patients had at least another infectious disease at the time of strongyloidiasis diagnosis, being Chagas disease (21.8%), latent tuberculosis infection (6%), and schistosomiasis (5%) the most prevalent ones (S1 Table).

Among the 833 immigrants, South America was the most frequent geographical area of origin, and mean time of residence in Spain was 6.8 (SD 5.4) years. The median duration of the trip in travelers and VFR-immigrants were 45 (IQR 19.7–227) days and 30 (IQR 30–60) days respectively, and Sub-Saharan Africa and South America were the most frequent geographical areas of travel destination (S2 Table).

After requesting extra information of the included patients to the participant centers, it was available from 1062 (85.3%) patients. Regarding *S. stercoralis* infection diagnosis, patients could be classified as follows: definite in 233 (21.9%) patients, probable in 648 (61%) patients, and possible in 181 (17.1%) patients. Proportion of positive results for each microbiological technique was: 17.4% for stool formalin-ethyl acetate sedimentation, 22.5% for *S. stercoralis* fecal culture, and 98.8% for *S. stercoralis* serology. These differences were higher when analyzing the efficiency of the microbiological techniques in the 561 patients in whom all three
techniques were performed (see Table 2). Nine hundred and sixty-eight patients received specific treatment for strongyloidiasis; 94 patients did not return to the centers after the diagnosis, and treatment could not be offered to them. Ivermectin was the drug of choice in 90.4% of the cases. From the 968 patients who received treatment, 784 (81%) had at least one follow-up control, with median time of follow-up of 6 (IQR 3–11) months. Treatment outcomes of these patients were as follows: 301 (38.4%) cure, 350 (44.6%) probable cure, 54 (6.9%) treatment
failure, and 79 (10.1%) with no enough information. From the 54 patients considered treatment failure, only in four cases was due to larvae detection after treatment: a 40-years-old man coming from Sudan (treated with 200mcg/day x 2 days, larvae detected after 4 months), a 28-years-old women coming from Bolivia (treated with 200mcg/day x 2 days, larvae detected after 6 months), a 35-years-old women coming from Equatorial Guinea (with HIV infection, treated with 200mcg/day x 2 days, larvae detected after 3 months), and a 43-years-old man coming from Bolivia (treated with albendazole 400mg/12 hours x 7 days, larvae detected after 3 months). Therefore, 83% of the patients achieved a successful outcome (cure or probable cure). More information regarding the treatment and follow-up results is summarized in Table 3 and S3 Table.

Table 1. Clinical and demographic characteristics of patients with strongyloidiasis in +REDIVI (2009–2017).

| Clinical and demographic characteristics | Number of patients (n = 1245) |
|----------------------------------------|-------------------------------|
| **Demographic information**            |                               |
| Age, years                             | 38.3 (SD 12.9, 2–88)          |
| Gender, male                           | 590 (47.4%)                   |
| **Epidemiological groups**             |                               |
| Immigrants                             | 833 (66.9%)                   |
| VFR-immigrants                         | 338 (27.1%)                   |
| Travelers                              | 74 (6%)                       |
| **Immunosuppressant conditions**       |                               |
| None                                   | 1165 (93.6%)                  |
| HIV infection                          | 55 (4.4%)                     |
| Immunosuppressive drugs                | 15 (1.2%)                     |
| Organ transplantation                  | 5 (0.4%)                      |
| Other                                  | 5 (0.4%)                      |
| **Main reason for consultation**       |                               |
| Laboratory test alteration             | 512 (41.1%)                   |
| Health screening                       | 491 (39.4%)                   |
| Gastrointestinal symptoms              | 105 (8.4%)                    |
| Cutaneous manifestations               | 105 (8.4%)                    |
| Fever                                  | 33 (2.7%)                     |
| Respiratory symptoms                   | 9 (0.7%)                      |
| Bone/muscle manifestations             | 8 (0.6%)                      |
| Cardiovascular symptoms                | 7 (0.6%)                      |
| Lymphadenopathy                        | 4 (0.3%)                      |
| Other                                  | 17 (1.4%)                     |
| **Clinical presentation**              |                               |
| Non-complicated strongyloidiasis       | 1240 (99.6%)                  |
| Hyperinfection syndrome                | 5 (0.4%)                      |

NOTE. Data are reported as number (%) of patients or mean (SD, range)

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When analyzing potential factors associated with having a definite diagnosis, after the multivariate analysis, the presence of eosinophilia was the most strongly associated factor, with an OR of 3.83 (95%CI 2.09–7.04). Presence of moderate/severe eosinophilia and male gender were also associated with confirmed diagnosis, with an OR of 1.57 (95%CI 1.15–2.14) and 1.69 (95%CI 1.25–2.29) respectively. Conversely, the immigrant group (compared with VFR-immigrants and travelers) had a negative association with definite diagnosis, with an OR of 0.64 (95%CI 0.47–0.87). More information is depicted in Table 4. Regarding factors associated to
achieving the cure criteria, treatment with ivermectin (compared with treatment with albendazole) was the most strongly associated factor, with an OR of 2.34 (95%CI 1.23–4.47). Male gender, presence of eosinophilia, and definite diagnosis had a negative association for achieving the cure criteria (see Table 5).

Table 2. Laboratorial and microbiological information at the time of diagnosis of patients with strongyloidiasis in +REDIVI with available extra information (2009–2017).

| Presence of eosinophilia | Positive microbiological techniques | Number of patients (n = 1062) | Number of patients (n = 561) |
|--------------------------|------------------------------------|------------------------------|------------------------------|
| Mild (&lt;1.0x10⁸ cells/L) | stool microscopic investigation     | 490/867 (56.5%)              | 52/561 (9.3%)                |
| Moderate (1.0–3.0x10⁸ cells/L) | S. stercoralis specific faecal culture | 356/867 (41.1%)          | 84/561 (15%)                  |
| Severe (&gt;3.0x10⁸ cells/L) | S. stercoralis serology           | 21/867 (2.4%)               | 554/561 (98.8%)               |
| HTLV-1 infection         | Patients with positive results in all techniques | 9 /224 (4%)                 | 40/561 (7.1%)                |

| S. stercoralis infection diagnosis | Patients with 2 positive results | Number of patients (n = 1062) | Number of patients (n = 561) |
|-----------------------------------|---------------------------------|------------------------------|------------------------------|
| Definite                          | stool microscopic investigation | 233/1062 (21.9%)             | 37/561 (6.6%)                |
| Probable                          | S. stercoralis serology         | 648/1062 (61%)               | 8/561 (1.5%)                 |
| Possible                          | Culture + microscopic investigation | 181/1062 (17.1%)       | 4/561 (0.7%)                 |

| Positive microbiological techniques | Patients with only 1 positive result | Number of patients (n = 1062) | Number of patients (n = 561) |
|------------------------------------|--------------------------------------|------------------------------|------------------------------|
| Stool microscopic investigation    | stool microscopic investigation      | 173/995 (17.4%)              | 469/561 (83.6%)               |
| S. stercoralis specific faecal culture | Culture                           | 145/643 (22.5%)            | 3/561 (0.5%)                  |
| S. stercoralis serology            | Microscopic investigation           | 935/946 (98.8%)             | 0/561 (0%)                    |

NOTE. Data are reported as number (%) of patients.

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Table 3. Treatment and follow-up information of patients with strongyloidiasis in +REDIVI with available extra information (2009–2017).

| Drug of choice (n = 968) | Duration of the treatment | Follow-up (n = 968) |
|-------------------------|---------------------------|---------------------|
| Ivermectin              | Duration for ivermectin (n = 884) | Number of follow-up controls |
| 150mcg/Kg/day po         | 94 (9.7%) 1 day 31 (3.5%) | No control 184 (19%)   |
| 200mcg/Kg/day po         | 780 (80.6%) 2 days 840 (95%) | One control 520 (53.7%) |
| 200mcg/Kg/day po + sc    | 1 (0.1%) 4 days 9 (1%) | Two controls 264 (27.3%) |
| Albendazole 400mg/12 hours po | 84 (8.7%) 7 days 3 (0.4%) | Mean time of follow-up, months 6 (IQR 3–11) |
| Ivermectin 200mcg/Kg/day po + albendazole 400mg/12 hours | 9 (0.9%) 20 days 1 (0.1%) | Treatment outcome (n = 784) |
| Duration for albendazole (n = 93) | Cure 301 (38.4%) |
| 3 day                   | 5 (5.4%) Probable cure 350 (44.6%) |
| 5 days                  | 34 (36.6%) Treatment failure 54 (6.9%) |
| 6 days                  | 1 (1.1%) No enough information 79 (10.1%) |
| 7 days                  | 48 (51.6%) |
| 8 days                  | 1 (1.1%) |
| 10 days                 | 2 (2.1%) |
| 15 days                 | 2 (2.1%) |

NOTE. Data are reported as number (%) of patients or median (IQR). Po, peroral; sc, subcutaneous.

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Imported strongyloidiasis is increasingly being diagnosed in non-endemic countries, especially in Spain, who receives an important number of immigrants coming from endemic areas, although information comes from short series of cases [7–11]. Here we present a cohort of 1245 patients with imported strongyloidiasis in Spain: most of them were asymptomatic immigrants coming from South America, 82.2% had eosinophilia at the time of diagnosis, 6.4% of

| Table 4. Factors associated to definite diagnosis in patients with strongyloidiasis in +REDIVI with available extra information (2009–2017). |
|---------------------------------------------------------------|
| **Overall (n = 1062)** | **Definite diagnosis** (n = 233) | **Probable diagnosis** (n = 829) | **Univariate analysis** (P value) | **Multivariate analysis** (OR, 95%CI, and P value) |
| Mean age at diagnosis, years | 37.9 (SD 13.0, 2–88) | 39.4 (SD 12.7, 2–88) | 37.5 (SD 13.1, 2–82) | 0.058 | OR 1.01 (0.99–1.02), p = 0.181 |
| Gender, male | 520/1062 (49%) | 136/233 (58.4%) | 384/829 (46.3%) | 0.001 | OR 1.69 (1.25–2.29), p = 0.001 |
| Immigrants | 699/1062 (65.8%) | 136/233 (58.4%) | 563/829 (67.9%) | 0.007 | OR 0.64 (0.47–0.87), p = 0.006 |
| Mean age at diagnosis, years | 6.4 (SD 5.1, 0–34) | 5.6 (SD 4.3, 0–21) | 6.5 (SD 5.2, 0–34) | 0.050 | - |
| Presence of immunosuppressant condition | 71/1062 (6.7%) | 13/233 (5.6%) | 58/829 (7%) | 0.444 | - |
| LT-1 co-infection (n = 224) | 9/224 (4%) | 3/55 (5.5%) | 6/169 (3.6%) | 0.692 | - |
| Presence of eosinophilia (n = 1055) | 867/1055 (82.2%) | 219/232 (94.4%) | 648/823 (78.7%) | <0.001 | OR 3.83 (2.09–7.04), p<0.001 |
| Moderate/Severe eosinophilia (n = 1055) | 377/1055 (35.7%) | 113/232 (48.5%) | 264/823 (32.1%) | <0.001 | OR 1.57 (1.15–2.14), p = 0.004 |
| Presence of symptoms | 190/1062 (17.9%) | 40/233 (17.2%) | 150/829 (18.1%) | 0.744 | - |

**NOTE.** Data are reported as number (%) of patients or mean (SD, range).

* Immigrant group compared to VFR-immigrants and travelers.

b Time of residence in Spain in immigrant patients.

c Patients in whom the main reason for consultation were not laboratory test alteration nor health screening.

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### Discussion

Imported strongyloidiasis is increasingly being diagnosed in non-endemic countries, especially in Spain, who receives an important number of immigrants coming from endemic areas, although information comes from short series of cases [7–11]. Here we present a cohort of 1245 patients with imported strongyloidiasis in Spain: most of them were asymptomatic immigrants coming from South America, 82.2% had eosinophilia at the time of diagnosis, 6.4% of

| Table 5. Factors associated to cure in patients with strongyloidiasis in +REDIVI with available extra information (2009–2017). |
|---------------------------------------------------------------|
| **Overall (n = 705)** | **Cure (n = 301)** | **Not cure (n = 404)** | **Univariate analysis** (P value) | **Multivariate analysis** (OR, 95%CI, and P value) |
| Mean age at diagnosis, years | 39.8 (SD 12.8, 2–88) | 40 (SD 12.5, 5–71) | 39.2 (SD 13.1, 2–88) | 0.132 | OR 1.01 (0.99–1.02), p = 0.132 |
| Gender, male | 344/705 (48.8%) | 130/301 (43.2%) | 214/404 (53%) | 0.010 | OR 0.64 (0.47–0.88), p = 0.007 |
| Presence of immunosuppressant condition | 50/705 (7.1%) | 27/301 (9%) | 23/404 (5.7%) | 0.094 | OR 1.37 (0.73–2.56), p = 0.314 |
| HTLV-1 co-infection (n = 152) | 7/152 (4.6%) | 4/105 (3.8%) | 3/47 (6.4%) | 0.677 | - |
| Presence of eosinophilia (n = 704) | 608/704 (86.4%) | 241/301 (80.1%) | 367/403 (91.1%) | <0.001 | OR 0.49 (0.30–0.78), p = 0.003 |
| Moderate/Severe eosinophilia (n = 704) | 268/704 (38.1%) | 107/301 (35.5%) | 161/403 (39.9%) | 0.234 | - |
| Definite diagnosis | 178/705 (25.2%) | 50/301 (16.6%) | 128/404 (31.7%) | <0.001 | OR 0.44 (0.30–0.64), p<0.001 |
| Presence of symptoms | 122/705 (17.3%) | 63/301 (20.9%) | 59/404 (14.6%) | 0.028 | OR 1.42 (0.73–2.56), p = 0.094 |
| Treatment with ivermectin (n = 698) | 644/698 (92.3%) | 285/299 (95.3%) | 359/399 (90%) | 0.009 | OR 2.34 (1.23–4.47), p = 0.009 |

**NOTE.** Data are reported as number (%) of patients or mean (SD, range).

* Patients with not enough information to establish the treatment outcome were withdrawn from the analysis.

b Patients in whom the main reason for consultation were not laboratory test alteration nor health screening.

c Patients only analyzed when treated with a single drug (ivermectin or albendazole).

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them had some kind of immunosuppressant condition, almost half of them had another infectious disease (being Chagas disease the most frequent one), serology allowed the diagnosis in most of the cases, and treatment with ivermectin was associated with better outcome.

It is important to note that, although the percentage of patients with eosinophilia was very high (more than 80%), the eosinophilia could be due to other causes, since there were patients co-infected with other helminthiases such as schistosomiasis, soil-transmitted helminths, or filarial nematodes. Hence, the percentage of patients with eosinophilia could be overestimated.

As in other imported strongyloidiasis series, most of the cases were diagnosed in immigrants and VFR-immigrants [7–11]. It is important to note that strongyloidiasis in VFR may have been acquired prior to residence in Spain rather than during the recent travel to their country of origin. Strongyloidiasis in travelers, although described, is much more infrequent, and restricted to high risk travelers, such as long-term travelers, humanitarian workers, and backpackers. In a study performed in The Netherlands where 679 asymptomatic long-term travelers to the tropics (median travel duration of 12 weeks) were post-travel screened for intestinal parasites using molecular techniques, only one (0.1%) traveler was diagnosed of strongyloidiasis [13]. In our study, travelers represented 6% of the total study population, which is higher than in other studies; it could be due to the epidemiological profile of travelers registered in +REDIVI, with high percentage of high risk travelers (missionaries and humanitarian workers). Among immigrants and VFR-immigrants, South America was the most frequent geographical area of origin, which is concordant with the general immigrant population in Spain, probably due to cultural and historic reasons. Spanish Tropical Medicine Units are highly sensitized in offering Chagas disease screening in South American population, especially those coming from Bolivia. This is the reason of the high proportion of patients coming from Bolivia in our study, and the high proportion of Chagas disease-strongyloidiasis co-infection. This co-infection has been recently found to increase the likelihood to detect *Trypanosoma cruzi* DNA in peripheral blood, which may reflect the potential immunomodulatory effects of *S. stercoralis* infection [14].

The percentage of severe clinical manifestations in our study (0.4%) could be underestimated, since most of the patients were referred to the centers for screening their health status or eosinophilia investigation. From the five patients with hyperinfection syndrome, four had some immunosuppressant risk factor (80%), whereas only 6% of patients without hyperinfection syndrome were immunosuppressed. Two of them were receiving corticosteroids, which was the most related risk factor found in a recent systematic review that included 244 cases of severe strongyloidiasis [6]. HTLV-1 infection increases the risk of *S. stercoralis* infection, the risk of developing a severe strongyloidiasis, and the risk of treatment failure [15]. In our study, 9 patients had a positive HTLV-1 serological test (one of them presenting a severe strongyloidiasis), although the test was only performed in 224 patients (21% of the study population).

There is currently no gold standard for *S. stercoralis* infection diagnosis, since the available techniques are not sensitive enough. Larvae detection in stool or other samples gives a definite diagnosis, but the scarce and irregular larvae excretion in chronically infected patients makes the diagnosis challenging. Although parasitological techniques have been improving (concentration techniques, Harada-Mori, Baermann, fecal culture), these techniques are not sensitive enough [16, 17]; in our study, the proportion of positive results in patients in whom all three techniques were performed was 9.3% for the formalin-ether technique and 15% for the fecal culture. The presence of eosinophilia was strongly associated with higher proportion of definite diagnosis, which may reflect a higher larvae excretion, possibly related to a higher parasite burden in these patients. Male gender was also associated with definite diagnosis. Both factors have been previously associated with *Strongyloides* infection in studies where the infection was defined by positive parasitological techniques [18, 19]. Conversely, being immigrant had a
negative association with definite diagnosis. It could be explained due to the chronic situation of the infection in this group of patients (they have leaved the endemic region years ago), and the fact that many of them have been diagnosed during a health screening, without presenting neither symptoms nor eosinophilia. In recent years, DNA detection-based techniques (such as PCR) have been developed, showing higher sensitivity than classical parasitological techniques [20].

Fortunately, different serological techniques have been developed during the last decades in order to increase the sensitivity in the strongyloidiasis diagnosis, being the ELISA-based techniques the ones with highest sensitivity [21]. Previous studies have shown the usefulness of *S. stercoralis* serology in the diagnosis of strongyloidiasis, especially in those presenting eosinophilia with negative coproparasitological examination [8, 22]. Serology allowed strongyloidiasis diagnosis in 78.1% of the patients in our study, becoming a key tool in the management of these patients. Moreover, serological tests are also useful to evaluate the treatment response, although a uniform criterion to define cure has not yet been established. In general, a negative serology or a 50–60% decrease in the optical density after treatment is considered a cure criteria or a successful outcome, which was the criteria used in the present study, although the time when this decrease should be determined is not well established [8, 23–25]. Nevertheless, *S. stercoralis* serology may overestimate the prevalence of strongyloidiasis, since it may remain positive after resolution of the infection or could cross-react with other helminth infections. It has been suggested that increasing the cut-off value for positive serology may increase the specificity of the serological test in the diagnosis of confirmed strongyloidiasis [26]. Other diagnostic strategy could include a combination of different techniques, such as classical parasitological methods, serology, and even PCR.

Some clinical trials have compared the efficacy between ivermectin and albendazole for the treatment of chronic strongyloidiasis, with a cure rate ranging from 76% to 98% and 38% to 78% respectively. All these clinical trials included a relative low number of subjects, used different treatment schemes and dosages, and assessed the treatment success using classical parasitological methods. As mentioned before, this strategy may have overestimated the cure response due to lack of sensitivity [27–32]. A recent Cochrane review has performed a meta-analysis comparing the efficacy of ivermectin versus albendazole including 478 participants, showing higher parasitological cure with ivermectin (RR 1.79, 95%CI 1.55–2.08) with moderate quality evidence [33]. Although our study had a retrospective design, a high number of patients (698 participants) could be evaluated to compare the efficacy of ivermectin versus albendazole; and more interestingly, our study included serological techniques during the follow-up, that increases the possibility to detect treatment failures. Even though the evidence of ivermectin superiority, almost 10% of the patients in our study were treated with albendazole; it could be explained because most of these cases took place at the beginning of the study (the evidence was less than nowadays), and the availability of ivermectin in some Spanish regions is limited. Patients with eosinophilia or with definite diagnosed were less likely to achieve the cure criteria; although it could reflect higher parasitic load in these patients, the retrospective nature of the study makes difficult the interpretation of these results. Male gender was also negatively associated to cure, which has been previously described in other studies [30].

The results of the previously mentioned clinical trials are very different to and contradict with the results of a recent observational study performed in a non-endemic area (Buenos Aires, Argentina). In this study, 21 patients with strongyloidiasis were treated with ivermectin; during follow-up, in 14 patients larvae were detected in stools by conventional methods, and *S. stercoralis* DNA was detected by PCR in all patients [34]. Although these results are surprising, they should be taken with caution, since come from a small observational study, and stronger evidence (coming from clinical trials) shows high efficacy of ivermectin. In our study, although
79 patients were considered treatment failure, only in 4 patients larvae were detected by conventional methods. Management of strongyloidiasis treatment failure is not standardized; the general attitude of the participant centres in our study in these situations was to repeat a course of ivermectin or to treat with a long course of albendazole.

This study has some limitations given its retrospective nature. Patients were diagnosed through different serological tests, and information regarding the number of stool samples tested in each patient was not collected, and it could influence in the sensitivity of fecal-based diagnostic techniques. Diagnostic tests and time of evaluation during follow-up were different depending on the centre. Moreover, the number of patients treated with albendazole was very low, and it difficult the interpretation of the efficacy comparison between the two drugs. Hence, the results have to be interpreted with caution. Despite these limitations, the present study has very strong points, such as the high number of included patients, the participation of 22 centers distributed through different Spanish regions, and reflects how Tropical Medicine specialists diagnose and manage patients with strongyloidiasis in a non-endemic setting under real conditions. The collaborative networks, such as the Spanish +REDIVI network, give the possibility to gather information from high number of patients, being very useful to study some rare manifestations from common diseases [35]. It will allow us to further investigate situations such as severe strongyloidiasis or strongyloidiasis in immunosuppressed patients.

In summary, we present a cohort of 1245 patients with imported strongyloidiasis in Spain. Patients were mostly immigrants, and the majority of them were coming from South America. Serological tests allowed the diagnosis in most of the cases, and detection of larvae in stool samples was associated with male gender and presence of eosinophilia. Ivermectin was associated with higher probability of cure compared with albendazole. Given the risk of developing a severe presentation, screening of *S. stercoralis* infection should be mandatory in patients coming from or had traveling to endemic areas, especially in those with immunosuppressant conditions [36]. Although parasitological techniques are the only that can confirm the diagnosis, to include serological techniques is very useful both at diagnosis and follow-up. Whenever possible, ivermectin must be the drug of choice. And finally, given the high number of immigrants and travelers coming from endemic areas that live in Spain, although unlikely under current hygienic conditions, reintroduction of the disease in our country could potentially occur, which would have important public health implications.

**Supporting information**

- **S1 Checklist.** STROBE checklist. (DOCX)
- **S1 Table.** Supplementary Tables S1. (DOC)
- **S2 Table.** Supplementary Table S2. (DOC)
- **S3 Table.** Supplementary Table S3. (DOC)
- **S1 Appendix.** Members of the +REDIVI study group. (DOC)

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**Author Contributions**

**Conceptualization:** Fernando Salvador, Begoña Treviño, Elena Sulleiro, Israel Molina, Rogelio López-Vélez, José Antonio Pérez-Molina.

**Data curation:** Fernando Salvador, Begoña Treviño, Sandra Chamorro-Tojeiro, Adrián Sánchez-Montalvá, Juan María Herrero-Martínez, Azucena Rodríguez-Guardado, Núria Serre-Delcor, Diego Torrús, Josune Goikoaiteza, Zuriñe Zubero, María Velasco, Elena Sulleiro, Israel Molina, Rogelio López-Vélez, José Antonio Pérez-Molina.

**Formal analysis:** Fernando Salvador, Sandra Chamorro-Tojeiro, Israel Molina, José Antonio Pérez-Molina.

**Investigation:** Fernando Salvador.

**Methodology:** Fernando Salvador, Elena Sulleiro, José Antonio Pérez-Molina.

**Supervision:** Fernando Salvador.

**Validation:** Adrián Sánchez-Montalvá.

**Writing – original draft:** José Antonio Pérez-Molina.

**Writing – review & editing:** José Antonio Pérez-Molina.

**Writing – original draft:** Fernando Salvador.

**Writing – review & editing:** Fernando Salvador, Begoña Treviño, Sandra Chamorro-Tojeiro, Adrián Sánchez-Montalvá, Juan María Herrero-Martínez, Azucena Rodríguez-Guardado, Núria Serre-Delcor, Diego Torrús, Josune Goikoaiteza, Zuriñe Zubero, María Velasco, Elena Sulleiro, Israel Molina, Rogelio López-Vélez.

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