Histoplasmosis in Children; HIV/AIDS Not a Major Driver

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Abstract: The classification of histoplasmosis as an AIDS-defining illness has largely attributed its occurrence in people to the presence of HIV/AIDS especially in Africa. Prior to the advent of the HIV/AIDS epidemic, several cases of histoplasmosis were documented both in the pediatric and adult populations. Our review revealed 1461 reported cases of pediatric histoplasmosis globally in the last eight decades (1939–2021). North America (n = 1231) had the highest number of cases, followed by South America (n = 135), Africa (n = 26) and Europe (n = 4). Histoplasmosis was much more common in the non-HIV pediatric population (n = 1418, 97.1%) compared to the HIV population. The non-HIV factors implicated were, childhood malignancies (n = 207), such as leukemias and lymphomas as well as their treatment, lung diseases (n = 7), environmental exposures and toxins (n = 224), autoimmune diseases (n = 12), organ transplants (n = 12), long-term steroid therapy (n = 3), the use of immunosuppressive drugs such as TNF-alpha inhibitors (n = 7) malnutrition (n = 12), histiocytosis (n = 3), Hyper immunoglobulin M and E syndromes (n = 15, 1.2%), pancytopenia (n = 26), diabetes mellitus (n = 1) and T-cell deficiency (n = 21). Paediatricians should always consider or rule out a diagnosis of histoplasmosis in children presenting with suggestive of the above clinical conditions.

Keywords: histoplasmosis; children; low medium countries; HIV; childhood malignancies; immunosuppressants

1. Introduction

Histoplasmosis is a systemic fungal infection occurring globally. It is endemic in the Ohio and Mississippi river valleys of the United States of America, India and Southeast Asia [1,2]. The clinical presentation of histoplasmosis mimics features seen in tuberculosis (TB) and may be misdiagnosed as such [3]. Amongst risk factors associated with disseminated histoplasmosis (DH), HIV infection is attributed as the greatest factor predisposing patients to DH in the adult population [4]. However, before the advent of HIV/AIDS, several cases of DH were reported [5]. Additionally, a review on the documented cases of histoplasmosis in Africa, from 1952 to 2017 by Oladele et al. revealed 37 cases of histoplasmosis in the pediatric age group, of which 33 (89.2%) were non-HIV patients [4]. In another study by Arango et al. on histoplasmosis: results of the Colombian National Survey, 1992–2008, 17 cases of histoplasmosis were reported within the pediatric age group, two were patients with AIDS, four had hematological cancers, three had environmental exposures, while eight had unknown risk factors [6]. In yet another study on childhood
histoplasmosis by Lopez et al. from 1984 to 2010, of the total 45 cases found, only four cases were HIV positive patients [7]. The other cases were attributed to malnutrition, tumors and environmental exposure [7]. Pakasa et al., in a case series study also revealed a high incidence (41.744.4%, 15/36) of histoplasmosis in non-HIV children within the age range of 3–16 years [8]. A large percentage of histoplasmosis in the pediatric age group most likely occurs in the non-HIV population, despite being described as an AIDS-defining illness. The purpose of this review is to identify factors other than HIV infection predisposing individuals in the pediatric age group to histoplasmosis and to raise awareness on pediatric histoplasmosis in the non-HIV population.

2. Mycology Summary

Histoplasmosis is caused by the dimorphic fungus, *Histoplasma capsulatum*. The two varieties pathogenic to humans are *H. capsulatum* var. *capsulatum* and *H. capsulatum* var. *duboisii*. The third variety, *H. capsulatum* var. *farciminosum*, is an equine pathogen [2]. However, a new taxonomic rearrangement of *H. capsulatum* proposed by Sepúlveda VE et al. showed that it is composed of at least four different cryptic species that differ genetically and also in virulence: *Histoplasma capsulatum* (Panama or H81 lineage), *Histoplasma mississippiense* (NAm 1), *Histoplasma ohiense* (NAm 2) and *Histoplasma var americana* (LAm A) [2]. Human infection occurs by inhalation of macroconidia and short hyphal fragments which are converted into yeast forms in the alveoli. Transmission can also be oral, causing infection in the intestines [9]. Transplacental, congenital and perinatal transmission of infections were also reported [10–14]. Histoplasmosis is primarily a pulmonary disease but can be disseminated in individuals at the extremes of ages due to their weak immune system, in people with advanced HIV/AIDS disease and secondary to persistent exposures to Histoplasma conidia [2,15]. An extrapulmonary presentation involving the skin may also occur and may progress to cause bone damage [3].

3. Materials and Methods

Literature searches for publications on histoplasmosis in children preceding February 2021 were performed using PubMed, Google Scholar, AJOL, Cochrane Library, Africa-Wide: NiPAD, CINAHL (accessed via EBSCO Host) databases and grey literature to identify all published papers regarding the topic. The main search comprised individual searches using detailed medical subject heading terms for histoplasmosis in children, including broad terms such as “case reports of histoplasmosis in children” and/or “diagnosis” and “management” of histoplasmosis in children. References in all relevant papers were also reviewed for additional publications (“snowballing”) on histoplasmosis in children that may not have been published in the searched databases. No language restriction was applied. Only reports with the patients’ country of origin identified were included. Publications reporting cases of histoplasmosis in individuals greater than 18 years of age were excluded from the review, as well as studies involving histoplasmosis in non-humans. Publications without abstracts were also excluded. The case definitions employed were based on an international consensus statement by the European Organization for Research and Treatment of Cancer/Invasive Fungal Infections Cooperative Group (EORTC/IFICG) and the Mycoses Study Group (MSG) [16]. Case reports with undefined HIV status were assumed to be negative while case reports with defined HIV status (that is positive or negative) were documented as such.

4. Results

Histoplasmosis in children is reported globally with high endemicity in Northern and Southern America [5,17]. Our literature search revealed a total of 1461 documented cases of histoplasmosis in children reported globally (Table 1) dating from 1945 to 2021. The number of cases were highest in North America (n = 1231), with 76.7% (944/1231) occurring in the USA. Very few cases (n = 4) in the pediatric population were reported in
Europe and were found in immigrants from endemic regions [18–20]. The number of documented cases from other climes were, South America (n = 135), Africa (n = 65) and Asia (n = 26), respectively. More cases occurred in the non-HIV population compared to the HIV population, globally. The percentages of non-HIV patients with pediatric histoplasmosis across the various regions was 97.1% (1418/1461) and North America (96.8%, 1192/1231), South America (99.3%, 134/135), Africa (98.5%, 64/65), Asia (96.2%, 25/26) within each region, respectively, (Table 1, Figure 1). Pediatric histoplasmosis in America and China was caused by *Histoplasma capsulatum var capsulatum* (Hcc), while in Africa, it was largely due to *Histoplasma capsulatum var duboisii* (Hcd) (88.7%, 47/53). Most cases of disseminated histoplasmosis occurred in persons with compromised immunity, Table 2 [21,22]. These infections are also common at the extremes of age, i.e., in infants and the elderly, although cases of disseminated histoplasmosis were reported in apparently normal individuals without the usual risk factors of extremes of age or some form of immunosuppression [23,24]. Childhood cancers such as leukemias and lymphomas as well as their treatment may predispose to histoplasmosis infection [21]. Underlying lung diseases, environmental exposures and toxins, autoimmune disease, long-term steroid therapy, use of immunosuppressive drugs such as TNF-alpha inhibitors may also predispose to histoplasmosis in children who are HIV-negative. Other less common conditions include histiocytosis, hyper immunoglobulin M and E syndromes and pancytopenia, Tables 1–3 [21,22]. DH was also documented in both premature and mature infants with variable outcomes [25–29].

![Figure 1](image-url)  
*Figure 1.** Global distribution of histoplasmosis in children.*
Table 1. Global distribution of Pediatric histoplasmosis.

| Country          | N  | -HIV | +HIV | Hcc Hcd | Factors Other than HIV (Number) | None | References |
|------------------|----|------|------|--------|---------------------------------|------|------------|
| **NORTH AMERICA** |    |      |      |        |                                 |      |            |
| Canada           | 1  | -    | 1    | 1      |                                 | -    | [30]       |
| * Colombia       | 73 | 69   | 4    | 25     | LD (1)                           | -    | [6,7,31–34]|
| * Costa Rica     | 84 | 84   | -    | -      | EE (44), TD (20), HIg (10)       | 10   | [35,36]    |
| Ecuador          | 38 | 14   | 24   | 37     | Hlg (1)                          | 13   | [37–39]    |
| Jamaica          | 1  | -    | 1    | 1      |                                 | -    | [40]       |
| * Mexico         | 89 | 85   | 4    | 15     | AIC (3)                          | 82   | [41–44]    |
| Trinidad         | 1  | -    | 1    | 1      |                                 | -    | [45]       |
| * USA            | 944| 940  | 4    | 296    |                                 | 521  | [17,21,22,25–116] |
| **Total**        | 1231| 1192| 39   | 376    |                                 | 525  | 668        |
| **SOUTH AMERICA**|    |      |      |        |                                 |      |            |
| * Argentina      | 50 | 50   | -    | 11     | EE (1)                           | 49   | [117–119]  |
| * Brazil         | 27 | 26   | 1    | 5      | EE (5), Leu/Can (2), OT (1), SLE/Steroids (1), DM (1) | 16   | [120–128]  |
| Guyana           | 1  | 1    | -    | -      | OT                               | -    | [129]      |
| * Venezuela      | 57 | 57   | -    | -      |                                 | 57   | [130]      |
| **Total**        | 135| 134  | 1    | 16     |                                 | 12   | 122        |
| **EUROPE**       |    |      |      |        |                                 |      |            |
| Germany          | 1  | 1    | -    | -      |                                 | 1    | [18]       |
| Portugal         | 1  | 1    | -    | 1      |                                 | 1    | [19]       |
| Spain            | 2  | 2    | -    | -      | Pan (2)                          | -    | [20]       |
| **Total**        | 4  | 4    | -    | 1      |                                 | 2    | 2          |
| **AFRICA**       |    |      |      |        |                                 |      |            |
| Burkinafaso      | 3  | 3    | -    | -      |                                 | 3    | [131,132]  |
| Cameroon         | 1  | 1    | -    | -      |                                 | 1    | [133]      |
| CAR              | 1  | 1    | -    | -      |                                 | 1    | [134]      |
| Chad             | 1  | 1    | -    | -      |                                 | 1    | [135]      |
| Congo            | 26 | 26   | -    | -      |                                 | 26   | [136–145]  |
| Cote d’voire    | 4  | 3    | 1    | -      |                                 | 3    | [146–148]  |
| Gambia           | 3  | 3    | -    | -      |                                 | 3    | [149]      |
| Guinea-Bissau    | 1  | 1    | -    | -      |                                 | 1    | [150]      |
| Kenya            | 2  | 2    | -    | -      |                                 | 2    | [151]      |
| Mali             | 1  | 1    | -    | -      | EE                               | -    | [152]      |
| Nigeria          | 13 | 13   | 2    | 7      | Leu/Can (2), Mal (1), OT (1)     | 9    | [153–161]  |
| South Africa     | 5  | 5    | -    | 2      |                                 | 4    | [23,162–165] |
Table 2. Percentages of non-HIV factors predisposing to pediatric histoplasmosis.

| Factors                                | Number of Cases | %     |
|----------------------------------------|-----------------|-------|
| Environmental exposure                 | 224             | 15.8  |
| Leukaemia/Cancers                      | 207             | 14.6  |
| Pancytopenia                           | 26              | 1.8   |
| T cell deficiency                      | 21              | 1.5   |
| Hyperimmunoglobulinemia                | 15              | 1.1   |
| Organ transplant                       | 12              | 0.8   |
| Malnutrition                           | 12              | 0.8   |
| Tumour necrosing factor inhibitors     | 7               | 0.5   |
| Lung diseases                          | 7               | 0.5   |
| Autoimmune conditions (SLE, Rheumatoid arthritis) | 11         | 0.8   |
| hemophagocytic lymphohistiocytosis     | 3               | 0.2   |
| Steroids                               | 3               | 0.14  |
| Interferon-gamma Receptor 1 Deficiency | 1               | 0.07  |
| Mesenteric cyst                         | 1               | 0.07  |
| Diabetes mellitus                      | 1               | 0.07  |
| **Total**                              | **551**         | **38.8%** |

% = number of cases/total number of non-HIV cases (1419).

Table 3. Some case reports on pediatric histoplasmosis associated with non-HIV factors.
| Authors          | Case Patients (n) | Age/Sex/Country | Non-HIV Factors | Clinical Findings                                                                 | Diagnostic Method                                                                 | Treatment and Clinical Course                                                                 |
|------------------|-------------------|-----------------|-----------------|----------------------------------------------------------------------------------|----------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------|
| Hess et al., 2017 | 1                 | 11-year-old/Male/USA | T-cell ALL fever, malaise, and weight loss | CT scan: mediastinal adenopathy. Bone marrow: pancytopenia, noncaseating granulomas. Urine and serum histoplasma antigen was positive. Histology of bone marrow as-tested after 3 months of therapy. |                                                                                     | AmB therapy. Histoplasma antigen in urine and serum was unde-                                |
| Tu et al., 1991  | 2                 | 3-year-old/Male/USA | hyper-IgM syndrome fever, nausea, vomiting, stomatitis of the gingiva, abdominal pain, fatigue, and irritability. | Disseminated histoplasmosis was diagnosed by bone marrow biopsy and culture of esophageal biopsy |                                                                                     | R                                                                                              |
| Yilmaz et al., 1995 | 3               | 3-year-old/Male/Turkey | Hyper-IgM syndrome | Recurrent pulmonary infections, ulcerated lesion of about 10 cm on the left side of the face | IgM, 1380 mg/dl. Chest radiograph showed bilateral infiltration and bronchiectasis. Biopsy of skin lesion revealed intracellular yeast cells | ketoconazole 10 mg/kg/day, with immunoglobulin therapy                                         |
| Hess et al., 2017 | 4                 | 14-year-old/Male/USA | Persistent fever and neutropenia Desmoplastic small round cell tumor of the abdomen | Chest CT demonstrated a solitary calcified lung nodule. Histoplasma serology became positive with increasing titers of histoplasmosis yeast by complement fixation from 1:16 to 1:32. Diagnosis was pulmonary histoplasmosis Urine histoplasma antigen test: positive. Fungal culture grew *Histoplasma capsulatum* from lymph node biopsy. Histoplasmosis antibodies to both yeast and mycelia returned elevated (1:1024 yeast and 1:16 mycelia). |                                                                                     | R                                                                                              |
| Tschudy et al., 2010 | 5              | 8-year-old/Male/USA | Crohn Disease/Infliximab therapy fevers, cough, anorexia, and night sweats. |                                                                                     | Itraconazole, Posaconazole, AmB were used for therapy                                      |                                                                                               |

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5. Discussion

5.1. Pulmonary Histoplasmosis and TB

Histoplasmosis may be misdiagnosed as TB due to the similarities in their clinical presentation [3]. Mosam et al., 2006 reported a case of disseminated histoplasmosis in an 11-year-old female African child who initially received anti-Koch’s regimen for two weeks despite negative Mantoux test and negative AFB [3]. Additionally, though rare in the literature, cooccurrence of histoplasmosis and TB was also documented in the pediatric age group [150]. Clinicians should investigate more with respect to histoplasmosis when seeing this cohort of patients.
5.2. Cancers

Pediatric cancers are probably the most common predisposing factor to histoplasmosis infection in non-HIV patients. The presentation may be localized, i.e., pulmonary disease, or may be disseminated histoplasmosis infection although the latter appears to be commoner [21,61]. Most children have an underlying hematologic malignancy (leukemia or lymphoma) and the diagnosis of histoplasmosis may be challenging due to the similarity in appearance of pulmonary histoplasmosis to lung metastases of primary malignancies [21]. A lung biopsy may be required to distinguish the two [22]. The impairment of T-cell function in malignancies is the pathogenetic mechanism for the increase in susceptibility to Histoplasmosis in pediatric cancer patients [35,36]. The use of chemotherapeutic agents resulting in bone marrow depression and consequently predisposing the children to histoplasmosis and other opportunistic infections is another pathogenetic explanation for the increased incidence [21]. A 14-year review of 57 pediatric cancer patients by Anderson revealed that there were 61 episodes of histoplasmosis in the group with disseminated histoplasmosis being more common [61]. Fever was the most common presenting symptom and no mortalities were attributable to histoplasmosis infection but cancer therapy was often delayed or modified due to the infection. Most patients initially received unnecessary antibiotic therapy and they were mostly non-neutropenic. Hess et al. found that urine antigen detection which was highly sensitive for disseminated disease in oncology patients lacked sensitivity in patients with acute pulmonary disease and if utilized solely may lead to missed diagnoses [21].

5.3. Lung Diseases

Pediatric histoplasmosis most frequently manifests as an acute pulmonary disease [57]. Pulmonary histoplasmosis is known to more frequently affect individuals with underlying lung disease such as emphysema and less commonly chronic bronchitis [151]. Pulmonary histoplasmosis was also reported in individuals previously diagnosed with or treated for tuberculosis [151]. Asthma, the commonest chronic lung disorder of childhood is thought to predispose to pulmonary histoplasmosis with possible pathogenetic mechanisms including alterations in local immunity of the respiratory tract and chronic use of corticosteroid treatments [181]. The differential diagnosis should be considered in children with asthma who present with recurrent or prolonged symptoms and who have been exposed to histoplasmosis [181].

5.4. Environmental Exposure

Most cases of histoplasmosis in healthy children result in a clinically inapparent infection. Immunocompromised hosts, infants and the elderly may present with clinical and often disseminated disease [113]. The size of the inoculum which is a function of the degree of exposure is also implicated in the tendency to develop clinical infections. Daubenton et al. reported a case of a seven-year-old immunocompetent child who lived on a poultry farm and developed disseminated histoplasmosis [23]. Although the relevant environmental exposures considered were exposure to bat caves, non-cave exposures were also reported [23]. For instance, Haselow et al. reported that 12 children between the age ranges of 4–19 years had contracted histoplasmosis after attending a family bonfire that used bamboo obtained from a large red blackbird roost [90]. Seven of the children were ill enough to require admission but there were no reported fatalities. The large “inoculating dose” and the age of the children were probably important factors in the spread of the disease in this case.

5.5. Malnutrition

Malnutrition is a major association in pediatric morbidity and mortality. The World Health Organization reported in 2020 that malnutrition was a contributing factor in 45% of all mortalities in under-five children worldwide [184]. Malnutrition is the primary
cause of immunodeficiency worldwide and young children and the elderly are most affected [185]. Malnutrition leads to defects in both innate and acquired immunity and sets off a vicious cycle with malnutrition leading to poor immunity and increased risk of infections/more severe infections which will then lead to poor intake, increased metabolic demand for nutrients leading to even more malnutrition. Defects in cell-mediated immunity caused by malnutrition also predispose to histoplasmosis. Mata-Essayag et al., in a retrospective review of 158 cases of histoplasmosis in Venezuela noted that 6 out of 42 children (14.3%) with histoplasmosis were also malnourished [130]. Of these, four (67%) died [130]. Lopez et al., in a study of 45 Colombian children with histoplasmosis found that malnutrition was the most important risk factor and was associated with being found in 37% of the patients [7]. In an earlier study by Jimenez et al., malnutrition was an association in over two-thirds of children with disseminated histoplasmosis. The mortality rate from the study was 61% [41].

5.6. Organ Transplant

Organ transplant recipients are susceptible to infections due to the chronic use of immunosuppressive therapy [92]. While viral infections are far more common in solid organ transplant recipients, fungal infections are usually more serious [50]. Studies have reported on occurrences of disseminated histoplasmosis in pediatric kidney transplant patients which are by far the commonest solid organ transplants performed in the pediatric age group [92,94]. Hemophagocytic lymphohistiocytosis has often been reported in association with histoplasmosis infection in kidney transplant recipients and this complicates the management and often worsens the prognosis [50,92,94]. A case series of six pediatric kidney transplant patients in an endemic region of the USA put the attack rate of histoplasmosis among kidney transplant patients at 6.9/100 cases. One-third of the cases occurred in the first year following the kidney transplant and there were no fatalities [94]. Most of the patients were treated with Amphotericin B and then transitioned to an azole, with 83% receiving chronic suppression therapy with Itraconazole to prevent future occurrences, such therapy was usually given indefinitely. The review suggested there may be a case for histoplasmosis prophylaxis for pediatric transplant recipients in endemic areas [94].

5.7. Autoimmune Diseases/Steroid Therapy

Pediatric autoimmune diseases such as Juvenile Rheumatoid Arthritis (JRA) are frequently treated with steroids and other immunosuppressive agents such as methotrexate. Because of the low doses used, opportunistic infections are not very common in association with autoimmune steroid or immunosuppressive therapy [102]. Hunstad et al. reported a case of a seven-year-old female with JRA who developed histoplasmosis while on low-dose methotrexate. She had also recently been weaned off prednisolone treatment [102]. The authors also noted that histoplasmosis and other opportunistic infections (OIs) were less common in JRA patients receiving low-dose methotrexate than in adult patients receiving the same for rheumatoid arthritis. The rarity of fungal opportunistic infections such as histoplasmosis has also been noted among pediatric patients with Systemic Lupus Erythematosus (SLE) with the first case report of histoplasmosis diagnosed in a child with SLE being as recent as 2012 [121]. The patient in question presented with abdominal distension, splenomegaly and multiple mesenteric lymph nodes. The patient died of septic shock while undergoing treatment. While these reports are rare, the increasing number of children being diagnosed with autoimmune disorders and on treatment with steroids and immunosuppressive agents makes a case for Pediatricians and Rheumatologists to be vigilant in identifying such cases so outcomes can be improved.
5.8. Pancytopenia/Defective Phagocytic Function

Pancytopenia and defective phagocytic function usually in association with Haemophagocytic Lymphohistiocytosis (HLH) were noted as predisposing factors to pediatric histoplasmosis with Russ et al. reporting a case of histoplasmosis and HLH in a seven-month-old infant with vomiting and failure to thrive [186]. The report noted that the child lacked homozygosity for HLH or had an underlying immunodeficiency, and suggested that alterations in T-cell function increased her risk for HLH following the infection with histoplasmosis, although it was unclear which condition occurred first [186]. Children with defective T- and B-cell function as well as a defective phagocytic function also present with fungal opportunistic infections such as disseminated histoplasmosis. Such reports are not common in the literature and are mostly from the developed world, possibly due to better diagnostic facilities. Hasliza et al. reported a case of disseminated histoplasmosis in a two-year-old previously healthy child. The workup during the illness identified gross leucopenia with defective T-cell and phagocyte function [179]. The T-cell and phagocyte function were all found to have normalized three months after the illness.

5.9. Tumor Necrosis Factors Inhibitors

Tumor necrosis factor-alpha (TNF-alpha) inhibitors such as adalimumab, infliximab and etanercept are commonly used in the treatment of pediatric inflammatory bowel disease [49,78]. They may also be used as second-line agents for auto-immune disorders of childhood such as juvenile rheumatoid arthritis (JRA) and psoriasis. However, inhibition of the TNF-alpha pathway results in a predisposition to severe infections including histoplasmosis, which is the commonest invasive fungal infection in patients on TNF-alpha therapy [78]. Dotson et al. reviewed five cases of histoplasmosis in children receiving either infliximab or adalimumab for Crohn’s disease and noted that early diagnosis with discontinuation of anti-TNF medication as well as close follow-ups were key to achieving good outcomes [49]. The study did not however recommend screening for histoplasma before commencing anti-TNF alpha drugs. Another report in an eight-year-old male with Crohn’s disease on Infliximab who presented with histoplasmosis and concurrent Pneumocystis pneumonia further raised the issue of pre-commencement screening for opportunistic infections in children receiving TNF-alpha inhibitors [49].

5.10. Syndromes (Hyperimmunoglobulin E Syndrome, Hyperimmunoglobulin M Syndrome)

Hyperimmunoglobulin E syndrome also known as Job syndrome and Hyperimmunoglobulin M syndrome are associated with defects in immune function specifically with defects in immunoglobulin levels, neutrophil chemotactic defects, defects in T-cell signaling and cytokine abnormalities [74,83]. Impaired Th 17 cell differentiation is an immune defect consistently found in STAT 3 Job syndrome mutation and this mutation impairs protection against invasive fungal infections such as histoplasmosis [74,83]. Case reports from pediatric patients have shown that both syndromes are associated with pulmonary, disseminated [74,83], and even cutaneous histoplasmosis infection [184].

6. Conclusions

This study has revealed the high occurrence rate of histoplasmosis in the non-HIV pediatric population, a disease commonly described as an AIDS-defining infection. We have also documented the underlying condition more commonly associated with this disease in children. Misdiagnosis of histoplasmosis as tuberculosis due to their similar pattern of presentation is also emphasized. Physicians managing children with risk factors highlighted in this study and presenting symptoms suggestive of TB should be screened to exclude or include histoplasmosis.

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D.D., R.O.O., I.A., and Z.P. conducted a critical review of the manuscript. All authors have read and agreed to the published version of the manuscript.

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