

Hyperinfection Syndrome Case report and review of Literature.

Vasant Nagvekar¹, Rachel², V. Ramasubramanian³, Thirunarayanma⁴, Nambi Senthur³, Ghafur³

¹Ex-Registrar, Department of Infectious diseases Apollo Hospital, Chennai, Tamil Nadu, India.

²Microbiology Technician, Apollo Hospital, Chennai, Tamil Nadu, India.

³ID Consultant, Apollo Hospital, Chennai, Tamil Nadu, India.

⁴HOD, Microbiology Department, Apollo Hospital, Chennai, Tamil Nadu, India.

Received: March 2018

Accepted: March 2018

Copyright: © the author(s), publisher. It is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

ABSTRACT

Hyperinfection syndrome is a fulminant gram negative septic shock particularly occurring in immunocompromised patients and is caused by the nematode helminth, *Strongyloides stercoralis*. Hereby we are addressing this case for the physicians in the endemic areas that they should be aware of the bizarre manifestations of the disease that can mimic other diseases leading to misdiagnosis and medical errors.

Keywords: acute respiratory failure; disseminated strongyloides; Gram-negative sepsis; *Strongyloides stercoralis*.

INTRODUCTION

Hyperinfection syndrome is caused by the nematode helminth, *Strongyloides stercoralis*. Strongyloidiasis may have varied manifestations from asymptomatic disease to a life threatening infection called as hyperinfection syndrome which carries a mortality of almost 61- 83%.^[1,2] Hyperinfection and disseminated disease occur during amplification of the autoinfective life cycle. Hyperinfection syndrome presents as a fulminant gram negative septic shock usually in immunocompromised patients more commonly in patients who are on long term steroids and who is chronically infected.^[3] Classically, the syndrome presents in a chronically infected person after immunosuppressive therapy is initiated for an underlying condition. Other risk factors for disseminated *Strongyloides* include immunosuppressive therapy, transplantation, hematologic malignant disease, human immunodeficiency virus, malnutrition and other T cell mediated disorders. High clinical suspicion is required to diagnose this syndrome.^[4]

This case illustrates about hyperinfection syndrome with strongyloides and importance of clinical suspicion of strongyloides in a case of severe gram negative septic shock or ARDS in a patient who is immunocompromised and is on steroids. The review also highlights the varied presentation of various

clinical manifestations of hyperinfection and the treatment options.

CASE REPORT

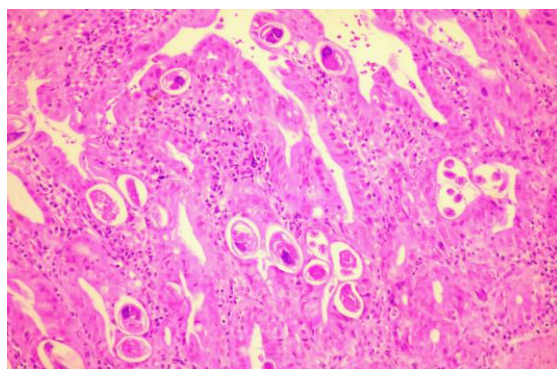
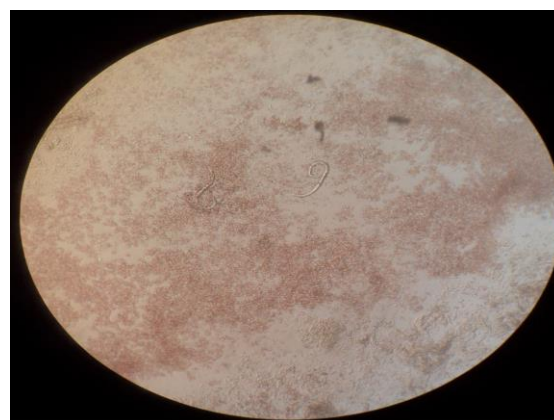
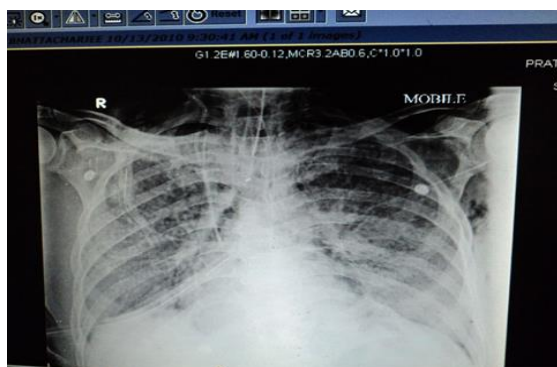
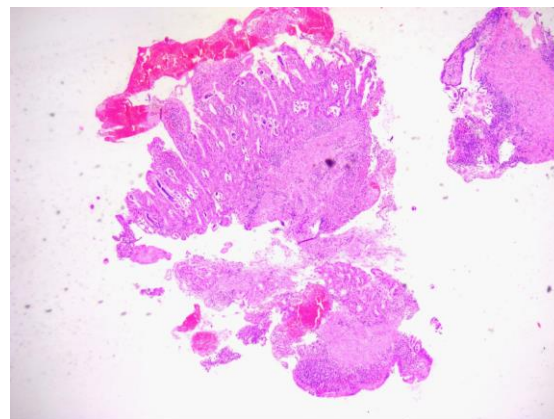
We report a case 6 months prior to the present admission. A 54 year male patient coming from Assam, north east India who had come for evaluation of anemia and was diagnosed to have autoimmune hemolytic anemia on Coombs test ++++ and after detailed investigations, including a bone marrow and ruling out all the causes of anaemia. He was started on prednisolone 60mg per day with monthly tapering doses by 10 mg after 2 months of prednisolone he developed a oral thrush and so his prednisolone was tapered to 25 mg per day in his local place at Assam and was treated with fluconazole. 3 months prior to this admission he started with weight loss, loss of appetite, loose motions off and on and progressive generalized weakness. 1 month prior to this admission he is admitted at Shillong in Assam with complaints of nausea, off and on greenish coloured vomiting, pain in abdomen, loss of weight by 10 kg. His investigations done then were Hb of 8.1gm%, white cell count of 8300 N76% and a platelet count of 3.05 lakhs. His Serum creatinine was 0.4mg%. His Serum Bilirubin was 1.2 mg% with a SGPT of 32 IU and his Total proteins were 3.3 with a S, Albumin of 1.6. He was treated symptomatically and with Ceftriaxone and he came for further work up at our hospital. He came to our hospital for haemetemesis and for his above complaints of three months duration. On admission he was conscious, afebrile with a pulse of 100/min, Blood pressure of 90/60 mmHG, a

Name & Address of Corresponding Author

Dr. Vasant Nagvekar
ID Consultant, Department of Infectious diseases
Apollo Hospital,
Chennai,
Tamil Nadu, India.

respiratory rate of 20/min. His CBC showed a Hb of 8.1gm%, WBC of 8600 N80% and platelets of 2.67lakhs. His Total proteins were 4.5 gm% with an albumin of 2.2gm%. His Serum creatinine was 0.8mg%. He was subjected for an Upper GI endoscopy which showed, Oesophagus and Stomach to be normal, Duodenum Second part showed nodularity, bleeding and unhealthy mucosa was reported as CMV? Lymphoma. Biopsy was taken and sent for histopathology examination. Meanwhile his Hb dropped to 4.6 gm% as he continued to have haematemesis and melena and was transfused with 3 units of packed red cells. Less than 24hrs of OGDScopy patient had hyperpyrexia with a temperature of 105 and further became haemodynamically unstable requiring multiple inotropes, had to be ventilated for desaturation. 2 sets of blood cultures were drawn and he was started on Meropenem and Vancomycin. His xray chest done showed an ARDS pattern.

subcutaneous ivermectin 12mg but died the next day. His blood cultures sent grew E.coli ESBL. This patient died due to a severe gram negative septic shock due to hyperinfection syndrome in a patient who was on long term steroid therapy. His duodenal biopsy showed larvae of strongyloides.



DISCUSSION

There was no obvious focus to explain his deterioration and since his xray showed ARDS like pattern with a background of history of steroid ingestion patient was subjected for a bronchoalveolar lavage. His investigations done then showed a Hb of 8.2gm%. a White cell count of 19090 N91% and a platelet count of 1.39 lakhs. His Coombs test was +++++. His Procalcitonin >10. He progressively deteriorated with increasing inotropes, and remained severely hypoxic with 100 Fio2. He became anuric was initiated on dialysis. Same day of Bronchoscopy a call from the microbiology department and demonstrated live larvae of strongyloides in the BAL sample. He was initiated on

Strongyloides the parasite is seen usually in the tropics and subtropics infecting about 100 million people in about 70 countries. It is endemic in Southern, Eastern, and Central Europe, Islands of the Caribbean, Latin America, Sub-Saharan Africa and Southeast Asia. It is very important that physicians should be aware of the endemic areas with strongyloides as failure to diagnose carries a high morbidity and mortality.^[1] Hyperinfection syndrome is not exactly defined, but the hallmark is an increase in the number of larvae in the stool and/or sputum along with manifestations confined to respiratory and gastrointestinal systems along with peritoneum.^[5] In Strongyloides hyperinfection syndrome clinical suspicion and diagnosis is often delayed upto a weeks time after hospital admission.^[6] In cases of hyperinfection the burden of larvae is so high that it can be seen easily in wet mounts of sputum or bronchoalveolar lavage fluid samples as in our case. In a person who has been chronically infected with strongyloides an immunosuppressed state causes enormous

multiplication and migration of infective larvae and is fatal. In presence of an immunosuppressed state there is a trigger and augmentation in the life cycle of the parasite. The triggering event in our patient going into gram negative septic shock after upper OGD scopy was, larvae present in the duodenum proliferated dramatically in the duodenum and migrated through the bowel wall, and then travelled through the venous system to the lungs and back to the small bowel along with the translocation of bacteria causing severe gram negative septic shock.^[3,7,8] The common organisms causing septic shock in hyperinfection are *Streptococcus bovis*, *Escherichia coli*, *Streptococcus fecalis*, *Klebsiella pneumoniae* or *Enterobacter* sp.^[3]

The manifestations of hyperinfection syndrome are divided, based on the system of origin, into intestinal and extraintestinal disease mainly involving the respiratory tract. The intestinal manifestations may present due to local larvae causing severe cramping abdominal pain, watery diarrhea, or malabsorption causing weight loss, nausea and vomiting and occasionally gastrointestinal bleeding due to intestinal erosions. Subacute intestinal obstruction can also be caused by strongyloidiasis and our patient had this symptom with bilious colour intermittent vomiting. In fact our patient had all the minor as well as severe intestinal manifestations of strongyloidosis.^[9]

The extraintestinal manifestations can present as Asthma-like symptoms such as cough and wheezing and it would be advisable in an endemic area to suspect hyperinfection syndrome in a patient who is on steroids or is immunocompromised. It can present as pneumonia and pulmonary hemorrhage with diffuse bilateral infiltrates on the chest x ray. Meningitis, Pericarditis, Myocarditis are rare manifestations. Gram negative Sepsis with ARDS due to autoinfection and gut translocation has already been described earlier.^[10,11] Patients receiving chronic steroids have an increased susceptibility to many different types of infections. The risk of infection is related to the dose of steroid and the duration of therapy. Although pyogenic bacteria are the most common pathogens, chronic steroid use increases the risk of infection with intracellular pathogens such as *Listeria*, many fungi, the herpes viruses, and certain parasites. Clinicians should consider both common and unusual opportunistic infections in patients receiving chronic steroids.^[12]

There is strong association of strongyloides and steroids and Siddiqui et al., have demonstrated the presence of steroid receptor on *Strongyloides stercoralis*, which could also play a role in the pathogenesis of hyperinfection syndrome and more systemic disseminated infection associated with corticosteroids.^[13] Corticosteroids, endogenous as well as exogenous, especially the exogenously administered have dual effect in causing

hyperinfection syndrome. It causes increase in ecdysteroid like substances which act as molting signals which causes an increase in the filariform larvae and on other hand steroids cause alteration in the mast cell function causing altered immunity and these dual together causes strongyloides hyperinfection syndrome.^[13,14]

Diagnosis: Due to the asymptomatic nature of the disease in its chronic stage diagnosis is difficult. Eosinophilia is an important laboratory finding in a patient with strongyloides but sometimes in hyperinfection you may have normal eosinophil count as in our case.^[15]

Microscopic examination of the stool and sputum may demonstrate larvae, Repeated stool samples require to be examined due to fluctuations in the excretion of larvae.^[16]

Enzyme-Linked Immuno-Sorbent Assay (ELISA) and Gelatin Particle Indirect Agglutination (GPIA) are antibody tests with sensitivities of 74.1% and 98.2% in a study by Human et al and with a specificity of 100%.^[17]

Treatment: The drug of choice for Strongyloides hyperinfection syndrome is Ivermectin 200 micrograms /kg daily till larvae are not seen at any site and repeat dose after 2 weeks. Intravenous and subcutaneous Ivermectin has been used in the same dose though it is not FDA approved for parenteral therapy. Alternative is albendazole 400mg twice a day or thiabendazole 25mg/kg twice a day till larvae are cleared and repeat dose after 2 weeks. Combination therapy with Ivermectin and Albendazole can be tried in severe cases.^[18,19]

Since these studies reported a daily dose of 200 g/kg/day, up to 14 consecutive days in the patient.^[20] (Ivermectin is now recognized as the drug of choice because it showed comparable and better rates of larval clearance than thiabendazole and albendazole, respectively, and fewer and comparable side effects than thiabendazole and albendazole, respectively.^[21]

CONCLUSION

Strongyloidiasis is a nematode infection which can present as a chronic asymptomatic to mildly symptomatic with gastrointestinal symptoms and at times can mimic as hyperreactive airway disease to fatal complications of hyperinfection syndrome and disseminated infection along with a host of other potential complications like gram-negative bacteremia and meningitis. Strongyloides is endemic in tropical and subtropical areas and it is very important for the clinician to know endemicity of the disease as it can present in a very severe form with a fatal outcome especially in people who are on steroids and immunocompromised states.⁽²²⁾ Diagnosis in both immune-intact and immunosuppressed individuals is usually by detection of larvae in sputum and stool samples and multiple stool samples are required due to the

fluctuations in excretion of larvae in stools.(23) Due to the asymptomatic nature of intestinal strongyloidiasis, and the risk for hyperinfection, screening of the population in endemic areas especially before considering immunosuppressive therapy is important. Physicians in the endemic areas should be aware of the bizarre manifestations of the disease that can mimic other diseases leading to misdiagnosis and medical errors.

REFERENCES

- Siddiqui AA, Berk SL. Diagnosis of Strongyloides stercoralis infection. Clin Infect Dis 2001; 33:1040–1047)
- Leelarasamee A, Nimmannit S, Na Nakorn S, et al. Disseminated strongyloidiasis: report of seven cases. Southeast Asian J Trop Med Public Health 1978; 9:539–542.
- Igra-Siegmán Y, Kapila R, Sen P, et al: Syndrome of hyperinfection with Strongyloides stercoralis. Respir Infect Dis 3:397, 1981.
- Raja S Vadlamudi1, David S Chi1,Guha Krishnaswamy1Intestinal strongyloidiasis and hyperinfection syndrome, Clinical and Molecular Allergy 2006, 4:8
- Keiser PB, Nutman TB: Strongyloides stercoralis in the Immunocompromised Population. Clin Microbiol Rev 2004 , 17:208-217. PubMed Abstract)
- DeVault GA Jr, King JW, Rohr MS, Landreneau MD, Brown ST 3rd,McDonald JC. Opportunistic infections with Strongyloides stercoralis in renal transplantation. Rev Infect Dis 1990; 12:653–71.)
- Mahmoud AA. Strongyloidiasis. Clin Infect Dis 1996; 23:949–52; quiz 53.
- G.C.COOK, Strongyloides stercoralis Hyperinfection Syndrome: How Often is it Missed? Quarterly Journal of Medicine, New Series 64, No. 244, pp. 625-629, August 1987
- Keiser PB, Nutman TB. Strongyloides stercoralis in the immunocompromised population. Clin Microbiol Rev 2004; 17:208–17.)
- E Chu, W L Whitlock and R A Dietrich Chest 1990;97:1475-1477 Pulmonary hyperinfection syndrome with Strongyloides stercoralis.
- Boulware, Brett R. Hendel-Paterson and Patricia F. Walker Ashley M. Newberry, David N. Williams, William M. Stauffer, David R.* Gram-Negative Sepsis Acute Respiratory Failure and Strongyloides CHEST 2005; 128:3681–368).
- Risk of infectious complications in patients taking glucocorticosteroids. - Stuck AE - Rev Infect Dis - 01-NOV-1989; 11(6): 954-63; Abstract.
- Siddiqui AA, Genta RM, Berk SL: Strongyloides stercoralis. In Infections of Gastrointestinal Tract. Volume 70. 2nd edition. Edited by: Blaser , Smith , Ravdin , Greenberg and Guerrant . Philadelphia, Lipponcott, Williams & Wilkins; 2002:1113-1126]
- Siddiqui AA, Berk SL, Genta RM: Strongyladiasis In Tropical Infections Diseases. In Philadelphia: Elsevier Edited by: Guerrant RL, Walker DH, Weller RF. 2005 , 1274-1285.
- Schulte C, Krebs B, Jelinek T, Nothdurft HD, von SF, Loscher T: Diagnostic significance of blood eosinophilia in returning travelers. Clin Infect Dis 2002 , 34:407-411)
- Uparanukraw P, Phongsri S, Morakote N: Fluctuations of larval excretion in Strongyloides stercoralis infection. Am J Trop Med Hyg 1999 , 60:967-973. Abstract.
- Huaman MC, Sato Y, Aguilar JL, Terashima A, Guerra H, Gotuzzo E, Kanbara H: Gelatin particle indirect agglutination and enzyme-linked immunosorbent assay for diagnosis of strongyloidiasis using Strongyloides venezuelensis antigen. Trans R Soc Trop Med Hyg 2003 , 97:535-538. PubMed Abstract.
- Strongyloidiasis in Transplant Patients Alison C. Roxby, Geoffrey S. Gottlieb, and Ajit P. Limaye Clinical Infectious Diseases 2009; 49:1411–23)
- Chiodini PL, Reid AJC, Wiselka MJ, Firmin R, Foweraker J, 2000. Parenteral ivermectin in Strongyloides hyperinfection.) Lancet 355: 43–44.
- JEROME PACANOWSKI, MARIE DOS SANTOS, et al : SUBCUTANEOUS IVERMECTIN AS A SAFE SALVAGE THERAPY IN STRONGYLOIDES STERCORALIS HYPERINFECTION SYNDROME: A CASE REPORT Am. J. Trop. Med. Hyg., 73(1), 2005, pp. 122–124)
- Marti H, Haji HJ, Savioloi L, Chwaya HM, Mgeni AF, Ameir JS, Hatz C, 1996. A comparative trial of a single-dose ivermectin versus three days of albendazole for treatment of Strongyloid stercoralis and other soil-transmitted helminth infections in children. Am J Trop Med Hyg 55: 477–481)
- Concha R, Harrington WJ, Rogers AI: Intestinal strongyloidiasis: recognition, management, and determinants of outcome. J Clin Gastroenterol 2005 , 39:203-211
- Gilles HM. Strongyloidiasis. In: Strickland GT, ed. Hunter's Tropical Medicine. 6th edn. Philadelphia: WB Saunders Co., 1984: 641-645.

How to cite this article: Nagvekar V, Rachel, Ramasubramanian V, Thirunarayanma, Senthur N, Ghafur. Hyperinfection Syndrome Case report and review of Literature. Ann. Int. Med. Den. Res. 2018; 4(3): ME36-ME39.

Source of Support: Nil, **Conflict of Interest:** None declared