Abdominal neoplasia with sarcomatoid features as the presenting illness of a patient with a newly diagnosed HIV infection and no AIDS-related disorders. Case report, clinical and diagnostic features, and literature discussion

INTRODUCTION

Even at the time of extensive availability of fully effective combination antiretroviral therapies (cART), missed or delayed diagnosis of HIV infection or full-blown AIDS remains proportionally frequent, as well as HIV disease presentations with multiple, concurrent opportunistic diseases [1-5], including both AIDS-defining and non-AIDS-defining disorders, which often make more and more difficult a prompt recognition and management by complicating the differential diagnostic workup.

In particular, due to the dysregulation of cancer controlling immune-mediated mechanisms persisting in HIV-infected subjects also despite a completely controlled HIV replication achieved by cART, both AIDS-related and AIDS-unrelated neoplasms remain frequent during recent years, as also noticed in the recent experience of our reference centre [1,6], in both adults and children. Despite the virological and immunological effectiveness of cART, malignancies on the whole continue to involve patients living with HIV due to their increased life expectancy, the continued exposure to many factors supporting cancer development, including environmental and lifestyle co-factors, as well as several co-infections often accompany.

Why we describe this case

This case underlines the importance of a careful monitoring of HIV patients, because, even in the cART era, they may present with rare and rapidly lethal solid tumors.

Abstract

We aim to describe a patient with an already advanced HIV infection disclosed for the first time during a complex diagnostic workup, which detected a gross abdominal mass attributable to a poorly differentiated mesenchymal cancer with sarcomatoid features which rapidly led our patient to death, in absence of other potential HIV-associated opportunistic diseases. Although extremely rare and rapidly lethal, our case report underscores the need of all caregivers who follow HIV-infected patients also in the cART era to maintain an elevated attention toward infrequent, unexpected, and clinically atypical solid tumors, in order to ensure a timely diagnosis and management when possible.

Keywords: Rare solid tumors; Sarcomatoid features; Newly diagnosed HIV infection; Case report; Differential diagnosis

Tumore addominale con caratteristiche sarcomatoidi come esordio di malattia in un paziente con nuova diagnosi di infezione da HIV e nessun disturbo correlato all’AIDS. Caso clinico, caratteristiche cliniche e diagnostiche e discussione della letteratura

CMI 2014; 8(4): 115-120
http://dx.doi.org/10.7175/cmi.v8i4.961
ing HIV infection itself (i.e. HBV, HCV, HPV, HSV, HHV-8, and EBV infections, among others), and a persisting immune system functional imbalance favoring the onset of tumors, in comparison with an epidemiologically matched sample of general population, which is not harboring HIV infection [2-5,7].

In particular, malignancies which are frequent in the general population often present with a clinical pathomorphism in our patient population, as in a case of atypical prostate cancer [8], and in some anecdotal presentations of Merkel cell carcinoma [9], and bladder and gastric cancer, respectively [10,11]. Very frequent associations with opportunistic and non-opportunistic disorders complicate even considerably the differential diagnostic pathway in multiple described case reports [1,2,6,7,12], in particular two cases of nasopharyngeal and rhinopharyngeal carcinoma [12].

Aim of our report is to describe a patient with an already advanced HIV infection disclosed for the first time during a complex diagnostic workup, which finally pointed out a gross abdominal mass attributable to a poorly differentiated neoplasia with sarcomatoid features which rapidly led our patient to death, in absence of other potential HIV-associated opportunistic infections and diseases.

**CASE REPORT**

A 36-year-old homosexual man with a medical history including only surgery for a pilonidal cyst 6 year before, and serological anti-HBV markers demonstrating a prior hepatitis B infection, during the last 6 months suffered from increasing anorexia and weight loss, followed by an irregular, mild fever not responsive to empiric broad-spectrum antibiotic treatments. When hospitalized in a General Hospital of our metropolitan area, the early instrumental examinations disclosed a parenchymal thickening of the lower left pulmonary lobe associated with a bilateral pleural effusion. Other imaging studies showed an enlarged spleen volume, diffuse lymphadenopathies along the main abdominal vessels, and a moderate peritoneal effusion, while no relevant abnormalities were detected with regard to liver and biliary tract. Posed on an empirical therapy with piperacillin-tazobactam, the patient was referred to our in-patient centre.

Upon admission, the laboratory examinations showed a marked leukopenia (total white blood cells = 1,570/µl), anemia (due to an hemoglobin level = 8.9 g/dl), low serum iron levels (26 µg/dl) with elevated serum ferritin (1,456 ng/ml), and especially a very advanced T-cell immunodeficiency, as sustained by a total CD4+ count of only 19 cells/µl (3% of overall T cell subset). All other blood laboratory examinations and urinalysis tested within normal limits, save an increased C-reactive protein (PCR) value (4.07 mg/dl).

Microbiological studies pointed out an elevated HIV viremia (112,181 HIV-RNA copies/ml of a wild-type HIV-1 virus), in absence of other active infections, which were carefully ruled out. In particular, repeated blood cultures for bacteria, fungi, and mycobacteria tested negative, as well as serum Parvovirus B19, Cytomegalovirus, and Epstein-Barr virus DNA search (only isolated anti-EBV IgG antibodies were present). Leishmania and Toxoplasma gondii serologies proved negative, as well as Clostridium difficile and Cryptosporidium search in the stools, stool cultures for other bacterial pathogens, urine antigens of Streptococcus pneumoniae and Legionella spp., and serum cryptococcal antigenemia and culture. With regard to major hepatitis viruses, HCV serology proved negative, while positive anti-HBs and anti-HBc antibodies represented the expected remnant of the previous documented HBV infection, in absence of dosable HBsAg serum levels. Ultrasonographic examination of the neck disclosed multiple, further lymphadenopathies of reactive origin.

During our hospitalization, the empiric antibiotic therapy was initially simplified with ceftriaxone plus clarithromycin, and the day after admission a potent antiretroviral combination therapy was immediately introduced (emtricitabine-tenofovir, plus darunavir 1200 mg/day plus ritonavir 200 mg/day), which proved well tolerated by our patient. After a 9-day hospitalization characterized by substantially stable general conditions and isolated mild fever and diarrhea (probably attributable to the recently discovered HIV infection and the severe underlying immunodeficiency), our patient was discharged, with treatment implemented with a chemoprophylaxis against Pneumocystis jiroveci and Toxoplasma gondii (performed with atovaquone), fungi (with fluconazole), and atypical mycobacteria (with clarithromycin, maintained after the previous, empiric ad-
ministration for the presumed respiratory infection).

Because of the re-appearance of irregular fever and an increased, bilateral pleural effusion, our patient was hospitalized again after a couple of weeks. An ultrasonography-guided thoracentesis of around 1000 ml of fluid neither allowed a diagnosis (all microbiological, mycobacterial, and neoplastic cell searches tested negative), nor a stabilization of the massive effusion, and was associated with increasing dyspnea, respiratory insufficiency, chest pain, diffuse peripheral edema, persisting anemia (requiring red blood cell transfusion), a worsening cachexia, and a rapid deterioration of general clinical conditions. As a consequence, our patient was moved to the Pulmonary Division, in order to try a surgical approach to the prominent pleural effusion.

A left trans-thoracic parietal pleural biopsy (with positioning a of trans-thoracic drainage), allowed to detect a first-degree empyema, in absence of isolated microorganisms at microscopy, culture, and molecular testing. Due to the concurrent, elevated fever, and increased serum ESR (erythrocyte sedimentation rate) and serum C-reactive protein levels, an empiric antibiotic therapy with piperacillin-tazobactam was introduced, associated to an empiric treatment against atypical mycobacteria, all showing no significant effects against irregular fever and the rapidly deteriorating general status. Quantiferon test proved negative. Subsequently, an elevated plasma HHV-8 viremia (64,000 viral copies/ml), required a treatment with full-dose intravenous acyclovir. A bone marrow and a liver biopsy were also performed, which allowed to exclude hematological malignancies and eventual, other opportunistic infections and disease localizations. A positron-emission tomography (PET) showed multiple hypercaptation sites at both lungs and pleura, a right axillary adenopathy, and a diffuse but non-specific abdominal hypercaptation.

A further worsening of respiratory and general conditions occurred in the next few days, with appearance of obtundation, lethargy, blurred vision, hypotension, abdominal distension, tendency to intestinal sub-occlusion, and a persisting, diffuse edema. All these complications rapidly led to death despite an intensive supportive care performed in an intensive care unit of our hospital, the adjunct of meropenem and linezolid (among antimicrobial agents), and the prosecution of antiretroviral therapy and that of the empiric anti-mycobacterial treatment.

The necropsy assessment showed the following macroscopic findings: a moderate edema at lower limbs, an abundant pleural effusion, a mild pericardic effusion, and a diffuse abdominal effusion (characterized by a clear, yellowish fluid). The respiratory tract showed an hyperemic and edematous laryngeal-tracheal-bronchial mucosa. Both lungs had a diffuse, increased consistency, a red-greyish color, and contained a frankly increased amount of foamy fluid. When examining the abdominal tract, the attention was immediately drawn on a voluminous grey-yellowish mass of myxoid appearance, which incorporated the intestinal loops and the entire colonic tract, with an extensive and infiltrating behavior. The liver showed a significantly increased size. The remaining major abdominal organs (including pancreas, kidneys, adrenal glands, and spleen) did not show relevant abnormalities, as well as all the main deep lymph node stations.

At microscopic examination, the abdominal mass was constituted by a poorly differentiated neoplasia with sarcomatoid features (Figure 1).

A further typization with all the available histopathological technique was not feasible at our centre, thus confirming the extremely non-differentiated features of this neoplasm. The concurrent, massive, irregular lobular necrosis of the liver was referable to a terminal, vascular compression due to the above-mentioned intrabdominal mass, which rapidly enlarged during the last days of life of our patient.
DISCUSSION

When approaching the differential diagnosis of rapidly growing abdominal masses in HIV-infected patients, tuberculosis and atypical mycobacteriosis in their broad spectrum of possible presentations remain the most frequent etiologies, starting from the pre-cART era until now [2,13]. But this diagnosis is neither obvious nor rapid; in our recent experience, an expansive abdominal mass leading to intestinal obstruction required a very cumbersome and prolonged workup in order to exclude all possible infectious and neoplastic etiologies, and was finally diagnosed and specifically treated with a 5-month delay after its clinical appearance, since only a positive culture for Mycobacterium avium-intracellulare became positive from a biopsy specimen, when all other clinical, microbiological, molecular biology, imaging, and histopathologic studies did not allow a disease identification in the meantime.

To add complication, a deep HIV-related immunodeficiency may lead to the first recognition of opportunistic infestations, including that due to a newly recognized cestode, which was initially responsible for a rapidly enlarging abdominal mass in a 1996 report [14]. On the other hand, the cART-related immune reconstitution syndrome has been also described as the supporting cause of a pseudo-tumoral abdominal mesenteric granulomatous mass, caused by an underlying mycobacterial infection plus inflammation and edema [15].

When focusing our attention on malignancies, AIDS-related ones remain frequent occurrences also in the cART era, with Kaposi’s sarcoma as the leading cancer with mesenchymal origin [3-5,16-18]. However, among HIV-infected children, Kaposi’s sarcoma and non-Hodgkin’s lymphoma declined in their frequency during the cART era in a more significant way, compared with the same AIDS-associated malignancies observed in adults.

In a recent review of case reports of sarcomas other than Kaposi’s sarcoma in the immunocompromised host (as a whole), Bhatia and coworkers identified 176 non-Kaposi’s sarcomas, 75 of them occurring in people with AIDS [18]. Leiomyosarcomas were the most frequently reported sarcomas according to histopathological assessments (101 cases), followed by angiosarcomas (23 episodes), and fibrohistiocytic tumors (17 cases). As already observed since the pre-cART era [19], and during the cART era too [3,5,17-20], smooth cell muscle neoplasms like leiomyosarcomas linked with immunodeficiency and a concurrent Epstein-Barr (EBV) infection, and often interested unusual body sites [19-21]; leiomyomas were also reported with increased frequency among HIV-infected patients, once again in conjunction with EBV infection [20]. Only one case of liposarcoma of the mediastinum has been reported in the international literature: in the year 1988 a 27-year-old man with a newly diagnosed HIV infection, who still had preserved peripheral T-cell subsets (with an absolute CD4⁺ count of 660 cells/µl) [22]. Histopathologic studies performed on the unresectable chest mass demonstrated a well differentiated liposarcoma of the de-differentiated subtype [22]. The Authors underlined the unusual localization and the young age at presentation of this liposarcoma, which were deemed to be

Key points

- Nowadays, despite a completely controlled HIV replication achieved by cART, both AIDS-related and AIDS-unrelated neoplasms remain frequently diagnosed
- Particular attention should be deserved to patients with a low CD4⁺ lymphocyte nadir, or a persistently impaired immune recovery, even during the cART era
- Malignancies with high frequency in the general population often present with a clinical pathomorphism in HIV-infected patients
- Starting from the pre-cART era until now, tuberculosis and atypical mycobacteriosis remain the most frequent etiologies when dealing with the differential diagnosis of rapidly growing abdominal masses in HIV-infected patients, but several other etiologies have been reported in the literature in this patient population
- It is mandatory to maintain an elevated attention level toward infrequent, unexpected, and clinically atypical solid tumors, in order to ensure a timely diagnosis, and make possible a more detailed workup, and management when feasible
It is mandatory to maintain an elevated attention level toward infrequent, unexpected, and clinically atypical solid tumors, in order to ensure a timely diagnosis, and make possible a more detailed workup, and management when possible.

**Key points**

- Malignancies with high frequency in the general population often present with a clinical pathomorphism in HIV-infected patients also in the cART era, due to a persistently impaired immune recovery, even during the cART era.
- AIDS-related and AIDS-unrelated neoplasms remain frequently diagnosed, especially in patients who were still unaware of their underlying retroviral infection.
- Although extremely rare and clinically untreatable, our case report underlines the need of health care personnel who follow HIV-infected patients also in the cART era, to maintain an elevated attention level toward infrequent, unexpected, and clinically atypical solid tumors, in order to ensure a timely diagnosis, and make possible a more detailed workup, and management when possible.

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