HIV medicine

Primary infection by type 1 human immunodeficiency virus: diagnosis and prognosis

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Despite preventive interventions, transmission of type 1 human immunodeficiency virus (HIV) has not been controlled and the incidence of HIV infection has been estimated to be 40,000 new cases per year in the US. These seroconversions are observed mainly among the young population, only 44% of which used condoms in 1991 compared to 56% in 1988. In this same study, the proportion of young intravenous drug users was 2.1% in 1991 vs. 0.5% in 1988. The risk of infection from unprotected sex and the growing consumption of intravenous drugs is worrisome, and these developments represent a serious failure of preventive campaigns.

More than half of infected persons will manifest clinical signs in the days following HIV infection. This initial stage of the disease was described for the first time in an isolated case in 1984, followed by a series of patients reported by Australian researchers in 1985. Recently, the first controlled trial in which zidovudine was administered at the time of primary infection has been published, with beneficial results. The diagnosis of patients in this early stage is more than ever a necessity. The objective of this article is to review the clinical presentation and biological disturbances occurring during primary HIV infection and their relationship to prognosis. The studies cited are those with the most rigorous methodologies associated with pertinent results and published in reviews of internal medicine and infectious diseases sourced through MEDLINE.

Epidemiology

The symptoms of primary infection have been described in all populations at risk of HIV infection: homosexual men, heterosexual men and women, intravenous drug users, recipients of contaminated blood products, recipients of organs from infected donors, and accidentally infected workers in the healthcare sector. Until now, no study has reported on different clinical features according to risk factors. In 95% of cases, there is at least one clinical sign. Nevertheless, in the presence of mild symptoms or in their absence, fever may pass completely unnoticed. Usually, the signs or symptoms of primary infection by HIV last for two to six weeks after contamination and persist, on average, for two weeks. These symptoms may become severe and require hospitalisation; 42% of patients were hospitalised in a recent Swiss study.

Clinical manifestations

The clinical manifestations are numerous but we will report the most frequent.

GENERAL MANIFESTATIONS (Box 1)
In the series reported by Kinloch et al., 87% of patients had fever whereas only 53% of those in the series of Pedersen et al. presented this symptom. The average temperature was 38.6°C, and eight out of 27 patients had a temperature over 39°C. Myalgia and arthralgia sometimes accompanied the fever. The mean duration of these symptoms was 16 days and 23 days. Persistence of symptoms for more than 14 days seemed to be related to a poor prognosis.

MUCOCUTANEOUS MANIFESTATIONS (Box 2)
A skin rash is present in more than 60% of cases (figure 1). These eruptions are erythematous, maculopapular, nonpruriginous, and usually symmetrical. They affect the face, trunk and sometimes the extremities, but generalised eruptions are rare. Vesicular lesions have also been reported.

Mucosal ulcerations are frequently reported: 43% in the series of Kinloch et al. They are localised in the oral cavity, sometimes associated with tonsillitis,
Figure 1  Severe abdominal skin rash during an acute HIV disease

Figure 2  Genital ulcer during an acute HIV disease

and on the genital organs (figure 2). In the same series, none of the patients infected by the parenteral route presented with mucosal ulcers whereas 52% of sexually infected subjects did. Gaines et al.\textsuperscript{12} found similar results in seven out of 20 homosexual men, indicating that ulcers may be the site of viral inoculation during sexual intercourse. Of patients presenting cutaneous lesions, 52% also manifest mucosal lesions, whereas 84% of subjects presenting mucosal lesions also manifest cutaneous lesions.\textsuperscript{8} The association of these lesions thus strongly suggests primary HIV infection because of the limited number of differential diagnoses.

**LYMPH NODE MANIFESTATIONS**

Adenopathies are present in more than 50% of cases.\textsuperscript{7,8} They are usually not painful, but may be generalised or localised.

**NEUROLOGICAL MANIFESTATIONS (Box 3)**

HIV has been isolated from cerebrospinal fluid in primary infection, indicating the early involvement of this system. The most frequent neurologic symptoms are headache, retro-orbital pains and photophobia. Depression, irritability and mood swings may also be observed, suggesting early involvement of affective  

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Box 3
functions, but depression is most often associated with announcement of the diagnosis. Other neurologic symptoms, such as meningitis or radiculopathy, are sometimes observed but they are less common.

PULMONARY MANIFESTATIONS
Pulmonary manifestations are not frequent. Patients may present with a dry cough, most often associated with nonspecific interstitial pneumonopathy.

GASTROINTESTINAL MANIFESTATIONS (Box 4)
Non-specific digestive symptoms such as anorexia, nausea and diarrhoea have been reported. The most suggestive signs of immune deficiency, such as oral and/or oesophageal candidiasis, have also been observed in association with severe transient immunosuppression. Oral candidiasis and oral hairy leukoplakia may appear relatively soon after seroconversion since the cumulative proportion of patients presenting these complications is 12% at one year and 30% at three years. Such observations are important because these two types of buccal lesions are associated with a higher risk of developing acquired immunodeficiency syndrome (AIDS).

Biological disturbances
Transient lymphopenia of helper T lymphocytes or CD4+ has been observed (figure 3), followed by lymphocytosis due to an elevation of cytotoxic T lymphocytes or CD8+. Thrombocytopenia is also a frequent anomaly. It is usually moderate, mostly about $145 \times 10^9/l$. An increase in the erythrocyte sedimentation rate has also been reported. The primary HIV infection is accompanied by activation of the immune and inflammatory system which results in an elevation of $\beta$-microglobulin, neopterin and $\alpha$-interferon. The number of CD8+ lymphocytes increases, while there is a decrease of helper CD4+ T lymphocytes with a persistent inversion of the CD4+/CD8+ ratio within the first year of HIV infection. The mean blood level of CD4+, measured six months before seroconversion, was $0.999 \times 10^9/l$ whereas it was a maximum of $0.349 \times 10^9/l$ one year after seroconversion.

Frequently, detection of HIV antibodies by the usual techniques (ELISA) is negative when symptoms are present, so that it is necessary to repeat the test after a few days or even weeks to confirm seroconversion. On the other hand, the symptoms coincide with significant viral load confirmed by a strong transient viraemia, an increase of the viral antigen p24 and an elevated titre of provirus in mononuclear cells from peripheral blood. In 76% to 90% of patients, p24 antigen is present between the 6th and 15th day after the first appearance of symptoms, then becomes undetectable between the 14th and 27th day. Quantification of viral genome circulating in plasma now permits the evaluation of viral load in the primary infection stage and is a good reflection of the quantity of circulating virus. The decrease of p24 antigen and other viral particles in blood is the result of the patient’s initial humoral immunologic response, of an increased number of cytotoxic CD8+ T lymphocytes and the sequestration of the virus in secondary lymphoid organs such as lymph nodes.

Prognosis (box 5)

AT THE TIME OF THE PRIMARY HIV INFECTION

Age
Age at the time of seroconversion seems to influence the AIDS incubation period since haemophilic children develop AIDS more slowly than adult haemophiliacs. An Italian study of intravenous drug users with seroconversion confirmed that an advanced age at the time of seroconversion was associated with a poor prognosis; these results were confirmed by cases infected by the sexual route.

Presence and duration of symptoms
A study of 48 patients with seroconversion (19 symptomatic and 29 asymptomatic) showed that evolution towards AIDS and/or a CD4+ level below $0.200 \times 10^9/l$ was faster in symptomatic patients. Nevertheless, arbitrary classification between symptomatic and asymptomatic patients is difficult since mild symptoms may sometimes be overlooked. It appears more realistic to classify patients according to a scale of severity or duration of symptoms rather than their presence or absence. According to Pedersen et al., the risk of developing AIDS three years after seroconversion is eight times
Summary/learning points

- primary HIV infection is symptomatic in nearly 70% of cases
- signs and symptoms are similar to mononucleosis but various clinical features may occur
- type, duration and severity of clinical features are related to HIV disease progression
- antiretroviral treatment at time of primary HIV infection seems to have a beneficial impact on prognosis

Box 6

greater for persons with symptomatic primary infection lasting more than 14 days. Progression to group 4 complications (according to the CDC definition\textsuperscript{25}) is also more rapid in this group (78% vs 10%). It thus appears that immunologic and viral factors associated with the progression of the disease play an important role at this stage. However, numerous questions are still unanswered: What is the influence of immunologic status of the patient before infection on the duration of symptoms? What is the impact of symptomatic treatment on the duration and severity of symptoms? What is the role of pre-existing genital mucosal ulcerations which may facilitate infection and permit penetration by a significant viral inoculum?

According to a Dutch study,\textsuperscript{23} the presence of fever and cutaneous eruptions at the time of seroconversion has been associated with a more rapid progression towards AIDS. This study confirmed the relationship between the severity of symptoms in the course of seroconversion but only two symptoms were analysed.

Viral strains
Roos et al\textsuperscript{24} undertook a prospective study of viral and immunologic parameters in 19 patients with seroconversion. Three patients were infected by viral strains inducing multinucleated cells or syncytia after cell culture. In these three cases, diminution of CD4\textsuperscript{+} was most rapid, and two of them presented AIDS at six and 19 months, respectively. In 15 patients out of 16 infected by strains not inducing the formation of syncytia, the CD4\textsuperscript{+} level was normal at the end of follow-up (average 391 days) and the patients remained asymptomatic. In two patients infected by a viral strain inducing syncytia, the same viral phenotype was found in the contaminating individual, confirming the possibility of transmission of these viral strains. Recently, the transmission of viral strains resistant to zidovudine was associated with more rapid progression of the disease.\textsuperscript{25}

Viral and immunologic markers
Schechter et al\textsuperscript{26} performed a prospective study of 18 subjects who developed AIDS in a cohort of homosexual patients in whom the date of seroconversion was known. Four months before seroconversion, these patients presented a decrease of CD4\textsuperscript{+}, a lower CD4\textsuperscript{+}/CD8\textsuperscript{+} ratio, a diminution of haemoglobin, an elevation of IgA, IgG and circulating immune complexes compared to 54 patients with seroconversion but who had not progressed to AIDS. In 84 subjects with seroconversion in the Multicenter AIDS-Cohort study, an American prospective study which included only homosexual patients, 18 developed AIDS within 36 months. Neither the level of p24 antigen nor the level of anti-p24 antibodies was associated with progression to AIDS.\textsuperscript{27} Patients who had developed AIDS by 36 months had significantly higher CD4\textsuperscript{+} levels (>0.400 x 10\textsuperscript{9}/l) at the time of seroconversion than those who did not evolve to AIDS. Lifson et al\textsuperscript{28} undertook a prospective study of the immunologic characteristics of 24 homosexual patients who did not progress to AIDS in nine years in comparison with patients who developed AIDS and HIV-negative patients. An elevated level of \(\beta\textsuperscript{2}\)-microglobulin and CD8\textsuperscript{+} at the time of seroconversion was associated with a favourable prognosis among those whose disease did not progress. In the study cited above,\textsuperscript{22} the absence of secretion of anti-HIV antibodies of the anti-p24 type and the presence of p24 antigen are independent predictors of a more rapid progression of the disease.

Concomitant infections
Do concomitant infections influence the evolution to AIDS? Cytomegalovirus and Epstein-Barr virus, which are common in the general population, themselves induce immunosuppression. It is thus possible that the pathogenic effects of each virus are potentiated. However, co-infection with other infectious agents such as hepatitis B or C virus, \textit{Treponema pallidum} or type 2 herpes simplex virus has not proven, up to now, to worsen the prognosis.

Impact of treatment of primary infection with antiretroviral agents
Trials on the use of zidovudine in the asymptomatic or slightly symptomatic stages have given contradictory results.\textsuperscript{29} Nevertheless, treatment of the primary infection stage has been shown to be warranted.\textsuperscript{30} For this reason, two controlled trials have been conducted, of which one is ongoing and the other has been published.\textsuperscript{6,31} Patients received either 500 mg/day of zidovudine or placebo for six months; 15 months after inclusion, the treated group presented significantly less opportunistic infections with a higher CD4\textsuperscript{+} count than the placebo group (values at the limit of significance). Prospective follow-up of these patients will allow long-term evaluation of zidovudine. However, early
prescription of antiretroviral treatment may lead to the rapid emergence of resistant viral strains, notably to zidovudine, which may then be transmitted from one patient to another. We ignore, for the moment, the question of whether contamination by this type of viral strain modifies the clinical presentation of the primary infection and whether the strain may again become sensitive to zidovudine in the absence of its administration.

PRIOR TO THE PRIMARY HIV INFECTION

Why do certain patients manifest symptomatic primary infection? Are there predisposing factors or a particular susceptibility to HIV infection? Little is known about this. The stage of the disease in the person who infects his or her partner appears to have an impact. Patients infected by the administration of contaminated blood products present symptomatic primary infection more frequently when donors develop AIDS within 29 months of the donation. Similarly, the recipients progress to a symptomatic stage more rapidly when the donor quickly develops AIDS. These observations coincide with studies on circulating viral load which demonstrated a larger quantity of virus in blood in the advanced stage of the disease. Nevertheless, recent investigations have shown that the quantity of circulating virus is important in the asymptomatic stages as well. The hypothesis of genetic predisposition linked, for example, to a certain type of major histocompatibility antigen (HLA) which induces rapid diminution of CD4+ is appealing, but the results remain contradictory, notably for the HLAs DR2, DR3 and DR5. The HLA genotypes A1, Cw7, B8 and DR3 meanwhile appear to be associated in a significant way with a more rapid decrease of CD4+, while the HLA phenotype DR2 seems to be frequently accompanied by the presence of p24 antigen in the blood.

Conclusion

The symptoms and signs of primary HIV infection are now better understood, and all physicians, whether they are general practitioners or specialists, should recall this diagnosis in persons presenting clinical manifestations of acute infectious disease and at risk of HIV infection. The search for viral load, p24 antigen, associated with antibodies, are essential, since most of the time, antibodies are undetectable at this stage of infection. Tests to detect antibodies should be repeated to confirm seroconversion. Biological investigations will nevertheless permit the exclusion of other possible diagnoses. The prescription of zidovudine or other antiretroviral medications at this stage has not proven their long-term efficacy against the development of AIDS and survival. Nevertheless, the results of a recent controlled clinical trial have demonstrated a beneficial effect of zidovudine on the CD4+ count with a reduction in the incidence of opportunistic infections after an average follow-up of 15 months.

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**Medical Anniversary**

**CLAUDE BERNARD, 12 July 1813**

Claude Bernard (1813–78) was born at St Julien in the wine-producing Beaujolais area of France, where his parents toiled in the vineyards. He qualified in medicine in Paris (1839) and became a professor at the Collège de France and Hôtel Dieu, succeeding professor of anatomy Magendie.

Claude wrote voluminously on biology and physiology, particularly on gastric juice in nutrition, curare, carbohydrate metabolism, the pancreas and the liver, and he isolated glycogen in 1857. Ill-health forced him to spend much time at his old home St Julien, where he had bought the manor house where his parents’ employer had lived. He became president of the Académie Française, and was the first French scientist to receive a national funeral in 1878. – *DG James*