HIV infection and psychiatric illness

B Owe-Larsson, L Säll, E Salamon, C Allgulander
Karolinska Institutet, Department of Clinical Neuroscience, Section of Psychiatry at Karolinska University Hospital Huddinge, Stockholm, Sweden

Abstract
Objective: To review the clinical features and current knowledge on the treatment of psychiatric symptoms and disorders in patients with human immunodeficiency virus (HIV) infection. Method: We searched the PubMed database combining HIV/AIDS with different keywords for psychiatric diagnoses and symptoms (e.g. depression, mania, anxiety, psychosis, dementia, substance abuse) and for psychopharmacological treatment. The years covered by these searches included 1980 to 2008. Results: Patients with HIV infection are at an increased risk of psychiatric illness. Major depressive disorder and subsyndromal depressive symptoms, as well as anxiety disorder and substance abuse are more prevalent among HIV infected individuals than among the general population. HIV-associated neurocognitive disorders (HAND) are common among HIV patients, and HIV-associated dementia (HAD) is a serious condition during the acquired immune deficiency syndrome (AIDS) stage of HIV disease. Secondary mania and psychosis might be the first clinical symptom of HIV dementia. The introduction of highly active anti-retroviral therapy (HAART) has resulted in significant decreases in morbidity and mortality for HIV infected patients. HAART has also decreased the incidence of HAD, but does not give complete protection from this condition. The utility of psychotropic medications in HIV patients has not been studied sufficiently as a basis for guidelines, and more controlled trials are needed. Conclusion: Psychiatric illness is common in HIV infected individuals, and underlines the importance for screening not only for cognitive impairment but also for co-morbid mental disease in HIV-positive patients. Further studies of the neuropsychiatric complications during HIV disease and the use of psychotropics under these circumstances are clearly needed. A better understanding of the pathogenesis of HAD is essential to identify additional therapeutic strategies for prevention and treatment of this neurodegenerative disease. Studies are also needed for optimizing effective utilization of antiretrovirals into the CNS. Mania and psychosis secondary to HAD may be used as an indicator to initiate HAART, irrespective of CD4 count. Further research on the utility of HAART in the treatment of such acute neuropsychiatric symptoms associated with HIV infection should be initiated.

Key words: HIV-associated neurocognitive disorders; HIV-associated dementia; AIDS; Psychosis; Secondary mania; HAART

Received: 23-05-2008
Accepted: 17-06-2008

Introduction
HIV is a ribonucleic acid (RNA) retrovirus that is the causative agent for AIDS. HIV belongs to the Lentivirus genus in the family Retroviridae, characterized by a replication cycle in which the viral RNA is reverse transcribed into a DNA proviral form that is integrated into the host cell genome. The retrovirus is composed of two copies of single-stranded RNA that codes for the virus’s 9 genes. The RNA is non-covalently linked to the core proteins, which in turn is surrounded by a viral envelope, which enables the virus to enter cells by binding to a specific cellular receptor located on the surface of cluster of differentiation 4 (CD4) cells.1 A few years after the discovery of HIV-1, a second virus, HIV-2, was found in West Africa. HIV-1 and HIV-2 differ regarding their content of the genome, resulting in lower AIDS rates in HIV-2 infected individuals.2 Once exposed to the body, the HIV virus antibodies may be detected after an acute flu-like illness, and then proceeds to infect specific cells of the immune system during an asymptomatic period that may last for years. In order to enter a cell, HIV must bind to CD4, typically found on T lymphocytes, blood monocytes, macrophages and some dendritic cells, and subsequently to one or a combination of several possible chemokine co-receptors, usually CCR5 and CXCR4.3 4 The late
stage of HIV infection, AIDS, begin after HIV has replicated itself into the T4 helper cells in the lymphatic system, over time causing a severely affected immune system. This stage is characterized by a generalized lymphadenopathy and AIDS related complex (ARC), leading to an immunosuppressed state with manifestation of a variety of opportunistic infections including cytomegalovirus (CMV), Epstein Barr virus (EBV), herpes simplex virus, John Cunningham (JC) virus, cryptococcosis, Pneumocystis carinii, toxoplasmosis and tuberculosis. In addition, patients may develop previously uncommon or unknown tumours, the most common of which are non-Hodgkin’s lymphomas and Kaposi’s sarcoma.

The existence of a specific infection of the CNS became apparent later and was confirmed by viral detection within the nervous tissue, together with the finding of multinucleated giant cells (MGC), a result of virus-induced fusion of infected macrophages. Characteristic neuropathological findings at autopsies in HAD are encephalitis with reactive astrogliosis, multinucleated giant cells, activated microglia, infiltration of mononuclear cells, myelin pallor and vacuolar myelopathy. These abnormalities are commonly seen within the central white matter, frontal cortices, basal ganglia, thalamus and brain stem. The myelopathy usually progresses in parallel with HAD, but occasionally develops before or without dementia, and manifests as a progressive spastic paraparesis with sensory ataxia. According to one model (the Trojan horse hypothesis), HIV enters into the CNS by migration of infected peripheral blood mononuclear cells through the blood brain barrier (BBB), and then resides primarily in perivascular macrophages and microglia. However, it is also possible that cell-free virus penetrates the brain by directly crossing the BBB. Various groups have found evidence of HIV infection also in astrocytes and oligodendrocytes, but neurons do not seem to be infected. However, it has been reported that neural progenitor cells are permissive to HIV virus. In macrophages/monocytes and microglial cells, HIV establishes a productive infection and is the source of transmission in CNS, while in other cells, as astrocytes, HIV only establishes a persistent, rather than productive infection.

HIV seems to penetrate into the CNS soon after the initial systemic infection in the periphery. However, there is little evidence that primary viral seeding result in a permanently productive infection within the CNS. During the period of immunodeficiency, i.e. AIDS, viral replication in the CNS is quite robust. The origin of this viral population is still unclear; it can be found. These findings represent increased water content and can be reversed with HAART. The severity of HIVE varies from mild cases to severe states with widespread inflammation and numerous giant cells. These variations could explain the observed range of clinical symptoms. The presence of HIVE shows a degree of correlation with HAD, but this is not an absolute association, and there are occasional cases with evidence of HIV at autopsy but no clinical history of dementia. The clinical features of HAD usually exhibit the hallmarks of a subcortical dementia, and are characterized by three key features: cognitive impairment, behavioural abnormalities and disturbed motor function.

**HIV-associated dementia (HAD)**

The involvement of the CNS was reported early in the AIDS epidemic. The concept of neuro-AIDS evolved between 1983 and 1988. Separate and overlapping diagnostic terms were created to define the psychiatric and neurological clinical symptoms related to HIV infection, collectively designated the AIDS dementia complex (ADC). HIV-associated dementia (HAD), HIV encephalitis (HIVE) is the pathological correlate of HAD, and should not be used to describe the clinical syndrome. HIV has a tendency to affect subcortical regions in the CNS. HIVE is particularly found in the basal ganglia and central white matter, but the neocortical grey matter and to a lesser extent the brainstem and cerebellum are sometimes involved, and HIVE may be present in any area of the brain. Magnetic resonance imaging (MR) typically demonstrates both cortical and central atrophy and corresponding ventricular enlargement. In addition, characteristic confluent signal abnormalities within the deep white matter can be found. These findings represent increased water content and can be reversed with HAART. The severity of HIVE varies from mild cases to severe states with widespread inflammation and numerous giant cells. These variations could explain the observed range of clinical symptoms. The presence of HIVE shows a degree of correlation with HAD, but this is not an absolute association, and there are occasional cases with evidence of HIV at autopsy but no clinical history of dementia. The clinical features of HAD usually exhibit the hallmarks of a subcortical dementia, and are characterized by three key features: cognitive impairment, behavioural abnormalities and disturbed motor function.

The initial cognitive symptoms often include memory loss, mental slowing, reading and comprehension difficulties and apathy. The typical mental and neuropsychiatric deficits of HAD are characterized by memory loss (selective for impaired retrieval), impaired ability to use acquired knowledge, personality changes (apathy, irritability, inertia), and general slowing of thought processes. Further neuropsychiatric sequelae in HIV-positive patients may include deficits in attention, concentration, language, executive skills, information processing, and subsequently aspects of motor functioning. This may also be accompanied by depressive withdrawal, impulsivity, manic symptoms, organic psychosis with paranoid features, sleep changes (hypersomnia, impaired judgement, disorientation, or even delirium.)
Considerable individual variability of presenting symptoms has been reported. HAD may be difficult to distinguish from depression in some cases, especially the initial symptoms which can be subtle and overlooked. Other conditions that may mimic HAD include opportunistic infections affecting the CNS and primary CNS lymphoma. CSF sampling and imaging studies should therefore be performed in the diagnosis of HAD to exclude opportunistic infections and other causes of cognitive impairment. Important tests of CSF include cell count and protein, culture (particularly mycobacterial and fungal), cryptococcal antigen, and polymerase chain reaction testing for toxoplasma, cytomegalovirus, Epstein Barr virus, John Cunningham virus and herpes virus.

As presented above, there might be significant cognitive deficits in cases of both asymptomatic HIV and AIDS, but these deficits are small in early stages of HIV and increase in the later phases of the illness. Motor functioning, executive skills, and information processing speed were among the cognitive domains that showed the greatest decline from early to later stages of HIV according to a meta-analysis made by Reger and colleagues. Early psychomotor slowing may predict the development of HIV-associated brain disease, and autopsy-verified HIV, and precedes clinical HAD by 1-2 years. However, the predictive value of quantitative motor test abnormalities for evolving cognitive impairment needs further delineation and testing. The severity of symptoms varies, however, in a study of 28 patients with early stage HIV infection (CD4 >280 cells/mm³), not receiving HAART, and followed during 7 years, no major deterioration in the neurological, psychological performance, neuropsychological or neuroimaging examinations could be revealed. Elevated HIV RNA in CSF may predict subsequent progression to neuropsychological impairment.

Amyotrophic lateral sclerosis (ALS)-like syndrome can occur in association with HIV infection, as well as sensory neuropathies, and neuromuscular syndromes as distal symmetric polyneuropathy (DSP), myopathy and HIV-associated neuromuscular weakness syndrome.

HAD follows a course of decreasing functional ability at a variable rate, resulting finally in a mute and paralyzed patient. The average survival with HAD is 6-9 months in untreated patients. HIV is probably the leading cause of dementia in people less than 40 years of age. HAD constitutes an independent risk factor for death due to AIDS.

In 1991, the Working Group of the American Academy of Neurology AIDS Task Force established diagnostic criteria for the diagnosis of HAD. According to these criteria, HAD requires a person to: 1) exhibit objective cognitive impairment, verified with standardized neuropsychological tests, in at least two domains (e.g., memory, attention, language, processing speed), 2) show impairment in performing activities of daily living (ADL), 3) experience either motor impairment or impaired emotional control or changes in social behaviour, 4) the pattern of cognitive impairment does not meet criteria for delirium (e.g., clouding of consciousness is not a prominent feature) and 5) exclusion of alternate neurological disorders through the use of laboratory studies, including CSF analysis when appropriate, and brain imaging by computed tomographic scanning or magnetic resonance imaging (MRI).

They also defined a less severe condition called minor cognitive motor disorder (MCMD). The essential features of MCMD were a history if impaired cognitive/behavioural function in two areas, causing mild impairment in work or ADL, does not meet criteria for HAD or HIV-associated myelopathy, and cannot be attributed to other etiologies. MCMD has been demonstrated to affect several important areas, including medication adherence and driving ability, and seems to have a high predictive value for subsequent detection of HIV at autopsy.

Some issues of the 1991 criteria may restrict their applicability. The number of domains of impairment that should be examined for diagnosis was not clearly defined; the degree of cognitive impairment was not fully specified; and there appeared to be some overlap between the criteria for mild HAD and MCMD. Finally, the classification did not admit mild forms of cognitive difficulties which had not developed to the point of interfering with everyday functioning. Therefore, an updated research nosology for HIV-associated neurocognitive disorders (HAND) was published, separating HAND into three categories:

1. HIV-associated asymptomatic neurocognitive impairment (ANI) is defined by 1) acquired impairment in cognitive functioning, involving at least two ability domains at least 1.0 standard deviation (SD) below the demographically corrected means (the neuropsychological assessment must survey at least the following abilities: verbal/language; attention/working memory; abstraction/executive; memory - learning and recall; speed of information processing; sensory-perceptual, motor skills), 2) the cognitive impairment does not interfere with everyday functioning, 3) the cognitive impairment does not meet criteria for delirium or dementia, and 4) there is no evidence of another pre-existing cause for the ANI.

2. HIV-1-associated mild neurocognitive disorder (MND), which is similar to the MCMD described in 1991, but more precisely defined: 1) an acquired mild-to-moderate impairment in cognitive function documented by a score of at least 1 SD below demographically corrected norms on test of at least two different cognitive domains, 2) the cognitive impairment interferes, at least mildly, with activities of daily living, 3) the impairment does not meet criteria for delirium or dementia, and 4) the impairment is not fully explained by co morbid conditions. Such a patient may show mild impairment in concentration, attention or memory, and for example complain of reading difficulties and being easily distracted due to poor concentration level.

3. HIV-1-associated dementia (HAD) requires according to these criteria 1) acquired moderate-to-severe cognitive impairment, documented by a score at least 2 SD below demographically corrected normative means in at least two different cognitive areas, 2) marked difficulty in ADLs due to the cognitive impairment, 3) the impairment does not meet criteria for delirium, and 4) the impairment is not adequately explained by co morbid conditions. Such a patient may for example show symptoms as delayed speech output, poor emotional and thought content with lack of spontaneity and social withdrawal.
Consequently, according to this nosology, the presence and degree of neurocognitive impairment constitute the fundamental criteria for diagnosis, while other criteria, e.g., motor disorders and emotional or personality changes are considered corroborative information. The criteria allow qualification of the progression of HAND according to activity. Determination of neurocognitive impairment should be based on appropriately normed tests. Different levels and examples of acquired impairment of activities of daily life (ADL) when diagnosing HAND is described by the authors. Before a diagnosis of ANI, MND or HAD is given, several co morbidity conditions has to be excluded in the patient. Such co morbid conditions include CNS opportunistic infections, medications with CNS effects, cerebral tumours, metabolic disturbances, or other developmental or acquired conditions unrelated to HIV disease.

A second level of co morbid condition is a contributing condition, i.e. the condition has had some substantive contribution to the neuropsychological impairment, for example a previous traumatic brain injury or a developmental disorder, but the effect of HIV is also considered to be significant. If a patient with suspected HAD also meet the criteria for a severe episode of major depression with significant functional limitations or psychotic features, or substance abuse, the diagnosis of HAD should be deferred to a subsequent examination performed at a time when the major depression has remitted or at least 1 month has elapsed since cessation of substance abuse. In the research nosology for HAND from 2007, it is further commented that the consensus was that even when major depression and HAD occur together, there is little evidence that pseudodementia exists and the cognitive deficits do not generally improve with treatment of depression.

In many resource-limited settings standardized neuropsychological examination procedures and other diagnostics tools, e.g computed tomography (CT) or magnetic resonance imaging (MRI) are not yet available. In particular, the neuropsychological tests and functional assessments may not have been validated in the language of many countries, or normative standards may be lacking. In these situations, the diagnostic guidelines for HAND from 2007 may be followed in principle, using clinical judgment and assessments aimed at establishing the same criteria.

If neuropsychological testing is not available, presence of cognitive impairment involving two or more ability domains may be detected by standardized mental status examinations, using appropriate demographic normative cut-offs if available. The HIV Dementia Scale (HDS) is a bedside screening tool to identify HAD and to monitor therapeutic effects. The HDS is comprised of four tasks that evaluate memory, psychomotor speed, construction ability and executive functions, and is specially designed for a condition with subcortical dementia. The HDS has a sensitivity of 80% and a specificity of 91%. It appears to be widely used because of its ability to distinguish patients with frank dementia from those who are cognitively normal. For the increasing number of patients with mild dementia and MND following the introduction of HAART, the HDS, as a screen, is not as accurate in detecting HAD as a more thorough neuropsychological examination. The International HIV Dementia Scale (IHDS) is another, new screening test. The anti-saccadic error subtest of the HDS has proven difficult for non-neurologists to administer, and the HDS also includes subtests (alphabet writing and cube-copying tests), which may be difficult for individuals with a non-Western educational background. The IHDS eliminates the antisaccades subtest and replaces the timed written alphabet and cube copy time subtests with tests of motor speed and psychomotor speed which can easily be performed across different cultures. The IHDS consists of three subtests: finger tapping, timed alternating hand sequence, and recall of four items at two minutes. In a study comparing patients in the U.S. and Uganda, its sensitivity was 80% and specificity 55-57%, making it useful as a screening instrument, both in the industrialized world and the developing world. As with the HDS, full neuropsychological testing should be performed to confirm a diagnosis of HAD. The Mini-Mental State Examination is an additional test that might be used to screen for cognitive impairment. Example of cognitive domains screened for in the tests presented above is given by Antinori and colleagues.

HAD has been estimated to occur in 15-30% of untreated HIV individuals and in approximately half of paediatric patients. AIDS patients in the late stages might have a more than 50% prevalence of HAD, and autopsy studies have revealed that up to 90% of patients with AIDS may have pathological indicators of HIV. HAD and myelopathy generally do not develop until advanced HIV infection. HAD may however, be the only presenting manifestation of AIDS, but is rare among otherwise healthy HIV-infected persons. Mild forms of cognitive impairment, MCMD, has been reported to exist in at least 30% of symptomatic HIV-positive adults. The combined measurement of plasma HIV RNA and CD4 lymphocytes predicts the prognosis of HIV/AIDS, and CSF levels of HIV RNA correlate with the severity of neurological deficits, at least before the era of HAART.

The WHO Neuropsychiatric AIDS study examined the prevalence of HIV-associated neurological and cognitive abnormalities in five geographical regions worldwide 1990-1991, using a neuropsychological test battery, a neurological examination, a structured interview for the diagnosis of HAD and a functional assessment. The prevalence of HAD for symptomatic HIV-positive patients was 4.4-6.9% in both Kenya and Democratic Republic of Congo, which was similar to the prevalence in Germany and Brazil. The prevalence of HAD in Africa was quite low in this study, possibly because the HIV epidemic was still in its early stages in this region in 1990-1991. In other studies from Africa, the frequencies of HAD in Northern Tanzania (1989) was seen in 54% of cases, in 8.7% of cases in Kinshasa, Democratic Republic of Congo (1992), and in 16% of cases in a rural Ugandan hospital (2002). However, formal neuropsychological testing was not performed as part of these studies, and differences in the diagnostic criteria for dementia could also account for the variance in the prevalence of HAD in these case series. Wong and colleagues made detailed sociodemographic, medical history, neurological, neuropsychological, and functional assessments on 78 people with HIV (mean CD4 cell count 219 cells/mm³) and 100 people who were HIV negative in Kampala, Uganda. HAD was present in 31% of all patients with HIV and an additional 47% of the patients met the criteria for mild cognitive impairment on neuropsychological testing.
Important risk factors for a diagnosis of HAD included subjective memory complaints, low education, low performance on the IHDS, advanced age and low CD4 cell count. Each additional 10 years of age conferred a greater than twofold risk of HAD. The study included HIV-positive individuals who had more advanced HIV infection than those in the WHO study. Salauw and colleagues studied the cognitive function in 60 asymptomatic, treatment-naïve HIV-positive Nigerians. These individuals differed significantly from individually matched, HIV-negative control subjects, and 56.6% showed cognitive impairment (at least one test -2 SD), compared to 13.3% in the control group. The CD4 cell count of the HIV-positive subjects had no significant correlation with the cognitive test scores, suggesting that cognitive changes during asymptomatic and early symptomatic HIV infection were not the result of immunosuppression in this group of patients.

The different frequency of HAD in different regions raise the possibility that HIV subtypes, or clades, might have different biological properties with respect to their capacity to cause HIV-associated cognitive impairment. In North America and Europe, clade B is the predominant HIV subtype, while in Africa several different subtypes with different distribution are known. The predominant HIV subtype in Ethiopia and Southern Africa, while D and A subtypes predominate in Uganda. In a recent study from Ethiopia, HIV-associated cognitive impairment was evaluated in 73 HIV-positive individuals (CD4 cell count 260 cells/mm³) and 87 HIV-negative individuals. The HIV-positive subjects had slower performance than the HIV-negative individuals in the finger-tapping test, but there were no differences between the performance of HIV-positive and HIV-negative individuals in any other neuropsychological test, or in the IHDS total score. However, Joska and colleagues that HAND was present in 23.5% in a sample of 536 HIV-patients (of which half were on antiretroviral medication) from South Africa. It has recently been shown in a mouse model, that HIV-1 clade B have greater impact on the induction of neuropathogenesis than clade C. If this has clinical significance in humans remains to be determined. The occurrence of neurocognitive impairment in HIV positive individuals during the era of HAART is given at the end of this review.

Psychiatric manifestations of HIV disease

The interplay between biologic, psychologic and social circumstances associated with HIV disease can be complicated. Careful diagnosis and treatment of psychiatric disorder in HIV positive patients are important as psychiatric manifestations can have serious effects if not identified. The occurrence of cognitive impairment in case of HAND might complicate a diagnosis of depression. Some psychiatric symptoms might also be directly related to HAD, as for example secondary mania or psychosis.

Knowledge of the metabolic pathways of psychopharmacologic agents as well as antiretrovirals is important because of potential adverse drug-drug interactions. Furthermore, patients infected with HIV have an increased sensitivity to side effects of psychotropics. Having this in mind, strategies similar to those that apply for the treatment of psychiatric disorders in the general population could be followed. However, controlled trials with HIV-positive individuals of several psychotropic agents are needed in the future. Given that psychiatric illnesses is present in almost half of HIV-positive individuals and that, of these patients, around 50% do not receive psychotropic medication, knowledge of the spectrum of mental disorders among HIV patients is vitally important.

Depression

Depressive spectrum disorders seem to be the most common psychiatric manifestations of HIV disease. Major depressive disorder (MDD) is more prevalent among HIV-infected individuals than in the general population, with estimated prevalence rates varying widely from 2% to 30%, or even up to 50% of the HIV-positive patients. Patients with HIV are 2-7 times more likely to meet diagnostic criteria for current MDD than individuals in the general population. The variable prevalence rates of MDD reported are probably explained by variations in study population and design, with differences in age, sex, education, ethnicity and stages of HIV/AIDS. For example, lower rates of MDD are found among patients who had not progressed to AIDS. Furthermore, different degrees of HIV cognitive impairment influence the patient's symptoms, which may be difficult to elucidate in the individual patient. Earlier prevalence studies have mainly focused on HIV-seropositive men. Estimates of depression among HIV-positive women range from 1.9% to 35% in clinical samples and from 30% to 60% in community samples. In a study of 93 HIV-positive women and 62 HIV-negative control women, MDD was significantly higher among the HIV-positive women (19.4%) than among the negative controls (4.8%).

MDD in HIV infected patients may be a primary consequence of CNS effects of HIV, a reaction to the stigmatization and emotional consequences of the diagnosis and coping with a serious medical illness, or a combination of these factors, thus constituting a heterogeneous group of affective disorder with neurovegetative confounding factors. Depression in individuals with HIV on HAART and without symptoms of HAD does not seem to be a clinically distinct subtype. Depression in HIV-positive patients might, however, influence the immune response. For example, MDD has been shown to alter the function of killer lymphocytes in HIV-infected women, resulting in increased levels of CD8 lymphocytes and HIV, and depression might lead to a progression of HIV disease and an increase in mortality rate. Depression might be unrecognized and untreated in many patients with HIV infection. Several of the cognitive symptoms of HAD may be difficult to differentiate from depression, and they may both precipitate similar symptoms as for example anorexia, insomnia, fatigue and pain, and they may both affect the aerotensregic system. The presence of anhedonia coupled with diminished mood in the morning indicate the presence of MDD. Depression may be the initial presentation of HAD, and may be associated with the progression of cognitive impairment. In addition, patients who demonstrate depression and cognitive impairment together may not respond well to treatment. Instruments to assess depression that allow one to separate cut items describing somatic symptoms from those concerning depressed mood should be used, as somatic symptoms associated with depression may also be caused by HIV infection. Some neuropsychological test may help in differentiating patients with HIV cognitive impairment with and
without depression. Depression does not seem to influence basal ganglia-mediated psychomotor speed, and such psychomotor evaluation of CNS function or dysfunction in HIV-positive patients might be used without having to take depression into account. Suicidality may be cause for psychiatric referral in those subjects who have been informed about a diagnosis of HIV/AIDS. In general, MDD in HIV patients lead to a decrease in adherence to HAART. However, appropriate psychiatric intervention may increase adherence to HAART, improve quality of life and decrease mortality. Therefore, detecting and treating MDD in HIV positive individuals is even more important.

Selective serotonin reuptake inhibitors (SSRIs) are effective in treating depression in HIV-infected patients. A more benign side effect profile with SSRIs has limited the use of tricyclic antidepressants (TCAs) to those HIV patients who do not respond to more modern antidepressants. There are a number of limitations of the available literature that has to be considered. In particular, the majority of controlled studies were conducted before the advent of HAART. Safety concerns have been raised concerning potential interactions between SSRIs and antiretroviral drugs. Protease inhibitors and non-nucleoside analogue reverse transcriptase inhibitors (NNRTIs) may affect the activity of cytochrome P450 isoenzymes, interacting with drugs metabolized by the same pathway including antidepressants, neuroleptics and anticonvulsants.

A serotoninergic syndrome in four patients receiving fluoxetine in combination with antiretrovirals that included ritonavir, efavirenz or sancquinar has been reported. To avoid such complications, a reduction of the initial SSRI dose, slow titration, and close monitoring for toxic reactions are recommended. Known interactions between antidepressants and antiretrovirals should always be considered before the medication is started.

The serotonin and noradrenaline reuptake inhibitors (SNRIs) represent an attractive alternative when treating depression in HIV-infected patients. Venlafaxine and duloxetine have been tried to a certain extent, but more studies are needed on the effectiveness and safety of these SNRIs when treating patients with HIV. These agents are also effective to treat chronic pain, which often is under treated in HIV individuals. Mirtazapine has been studied in a 12-week open-label study in 12 HIV-positive patients diagnosed with major depression. Mirtazapine was well tolerated and effective, and the participants reported a significant reduction of heaviness, loss of energy and fatigability, but larger studies should be accomplished. Also bupropion and reboxetine has been studied in small, open trials with HIV-positive patients, and accomplished. Also bupropion and reboxetine has been studied in a 12-week open-label study in 12 HIV-positive patients diagnosed with major depression.

Anxiety syndromes, adjustment disorder and posttraumatic stress disorder

Anxiety syndromes and symptoms are common among HIV-positive individuals, and often related to HIV symptoms, fatigue and physical limitations. The presence of persistent focus on death and mortality issues has been found to be common among HIV-positive patients in the symptomatic stage of the disease. The prevalence of generalized anxiety disorder (GAD) is increased in HIV-positive patients compared to HIV-negative individuals. Adjustment disorder, arising in response to difficult life events, is also common among HIV patients. Adjustment disorder was reported in around one-third of the patients in one study, and was prevalent at both asymptomatic and symptomatic stages of the HIV infection.

Posttraumatic stress disorder (PTSD) is another potential consequence of receiving a HIV/AIDS diagnosis. In one study, one-third of 61 HIV-positive patients met the criteria for a syndrome of PTSD in response to HIV diagnosis. PTSD in these patients was significantly associated with a pre-HIV history of PTSD from other causes, and with major depression arising after HIV infection. In a study from South Africa, the prevalence of PTSD was 30.6% in a sample of patients attending HIV clinics, reflecting the high rates of trauma and violence in the community according to the authors. PTSD was here significantly associated with HAND and the effect was dependent on the educational level.

Controlled studies assessing the efficacy and safety of anxiolytic medication in HIV-infected persons are lacking, including studies evaluating the efficacy of antidepressants for treating anxiety disorders in HIV-positive patients. In USA, benzodiazepines account for over 60% of all anxiolytics prescribed. Interactions between benzodiazepines and antiretrovirals have been reported. For example, due to its inhibitory effects on the CYP3 isoenzyme, low doses of ritonavir can significantly impair the clearance of alprazolam.

Mania

Mania is seen in heightened rates among individuals with HIV/AIDS, especially with the progression of HIV infection. Hypomanic or manic behaviour, including increased sexual activity and drug use, is an additional risk factor for contracting and transmitting HIV. Mania in a patient with HIV/AIDS may occur as part of a coexisting bipolar disorder, or it may be secondary to the effects of HIV infection on the CNS, treatments for HIV infection, or HIV-related secondary infections of the brain. Of these, HIV neurotoxicity has been postulated to be the most important factor in the pathogenesis of secondary mania among HIV positive individuals. In one study, patients with secondary mania (without family or personal history of mood disorder) and HIV presented later in the course of HIV infection, and were more likely to have developed AIDS, to have a low CD4 count and to have a higher prevalence of co morbid dementia or cognitive slowing compared to HIV-positive patients with primary mania, suggesting that secondary mania may be a direct effect of HIV on the CNS. The authors reported that the 17-month prevalence of mania in the studied population of HIV-positive patients was 1.4%, and the rate in the patients with AIDS was 8%. Patients with secondary mania had more symptoms and were more likely to show irritability, and less likely to display increased talkativeness. In another study, the prevalence of secondary mania over a 29 months study was 1.2% for HIV-positive patients, and 4.3% for those with AIDS. The patients in these studies were mostly males. Evidence for an etiological association between HIV disease and secondary mania was indicated in a study of HIV-positive patients with and without mania that demonstrated a protective effect from zidovudine, an antiretroviral agent penetrating the CNS. However, conclusions are tentative due to the small sample sizes in these studies. Secondary mania has been reported to occur...
with various disorders or processes that disrupts brain structure or physiology."

In a larger study from Uganda, 64 HIV-negative patients with primary mania were compared to 61 patients with HIV-related secondary mania. Compared to the patients with primary mania, the patients with secondary mania had more manic symptoms; they were more irritable, more aggressive and disruptive, more talkative, more likely to have a decreased need for sleep, and had higher rates of psychotic symptoms such as paranoid delusions (92% compared to 80%), visual hallucinations (93% compared to 16%) and auditory hallucinations (67% compared to 16%). Furthermore, they were more cognitively impaired (84% compared to 45% in primary mania), less educated, and more likely to be female. Among the patients with secondary mania, only half had been aware of their HIV status prior to their psychiatric hospitalization. In several of the patients, mania was the first revealing symptom of HIV infection. The majority of the HIV-positive patients (90%) did not have HIV-related illnesses other than the mania, lacking for example HIV wasting syndrome, Kaposi's sarcoma or opportunistic infections as Pneumocystis carinii or cryptococcal meningitis. According to the WHO clinical system for HIV/AIDS, 48% of the HIV positive patients were in stage 1 or 2, and 52% in stage 3 to 4. The authors suggest that secondary mania may be used as an indicator to initiate HAART.

Mood stabilizers as lithium and valproic acid (VPA) has been evaluated in pilot studies as adjunct therapy to ART (see below). They seem to be well tolerated and they might even improve neuropsychological performance. Consequently, they might also be used as mood stabilizers for treating bipolar symptoms in HIV-positive patients, but larger studies should be performed. For example, in one other study, a small case series of 10 HIV-positive patients, lithium was not well tolerated. Valproic acid should probably be used in conjunction with antiretroviral therapy, as VPA might increase HIV replication. Lamotrigine was tested in a randomized, placebo-controlled study and was effective in treating neuropsychiatric pain in 92 HIV-positive patients receiving antiretroviral therapy. However, the antiretroviral drugs lopinavir and ritonavir decrease lamotrigine plasma levels. Carbamazepine is an inducer of CYP3A enzymes and may increase the metabolism of the antiretroviral drugs lopinavir and delavirdine. On the other hand, ritonavir may raise the serum levels of carbamazepine. Further studies on the use of mood stabilizers in HIV-infected individuals are clearly needed, including those considering drug interactions.

**Psychosis**

Psychotic disorders can be classified into primary (e.g. schizophrenia, schizoaffective disorder) or secondary (e.g. psychosis caused by a medical condition such as HIV infection) disorders in the DSM-IV nosology. Thus, psychotic symptoms may be seen in HIV infected individuals due to several factors. Psychotic disorders may precede HIV infection. Patients with schizophrenia are at increased risk for co morbidity HIV infection because of an elevated likelihood of poor impulse control, impaired judgement, substance abuse and high-risk sexual behaviour. Although being sexually active tends to be less common among patients with schizophrenia, patients with schizophrenia who are sexually active are more likely to be engaged in high-risk behaviour. HIV disease tends to lead to greater morbidity and mortality in patients with schizophrenia than in the general public due to several reasons, such as difficulty complying with medical care and trouble explaining symptoms to medical personnel.

It has been shown that the HIV infection per se may be associated with psychotic symptoms. Studies have showed that new-onset psychosis in HIV positive patients occurred in 0.2% to 15% of the patients, with the highest incidence reported among patients in later stages of HIV disease and with HAART, suggesting that psychosis may be a direct effect of HIV infection on the CNS. 15% of 46 patients with HAD reported psychotic symptoms in one study, and these data were supported by a case study in which patients with psychosis showed greater neurocognitive impairment than HIV-positive patients without psychosis. The presenting symptoms are highly variable, but persecutory, grandiose, and somatic delusions are the most common symptoms, followed by auditory and visual hallucinations and affective disturbances. In a case study of 20 HIV-positive men with new-onset psychosis, all 20 presented with delusions, 12 with auditory and 9 with visual hallucinations, while mood symptoms occurred in 13 patients. Compared to schizophrenia, bizarre or complex delusions are less common in new-onset psychosis in HIV-infected individuals, and visual hallucinations and remission of psychosis are commonly found. Furthermore, as presented above, psychotic symptoms are commonly found among patients with secondary mania due to HIV disease.

A number of hypotheses have been proposed to explain the pathogenesis of new-onset psychosis in HIV disease: subcortical neurodegeneration caused by HIV itself or in the presence of other viral infections; psychosis secondary to HIV; brain damage from some other opportunistic infection, or an underlying dementia. The following factors have been reported to be associated with the development of psychosis in HIV infected patients: untreated HIV infection, cognitive impairment, dementia, and a history of psychiatric disease or substance abuse.

The use of HAART in HIV-infected patients may be expected to reduce the risk of new-onset psychosis secondary to HAD. HIV-positive patients with neuro-AIDS are more sensitive to side effects of antipsychotic agents, especially extrapyramidal side-effects (EPS), due to a loss of dopaminergic neurons. Modern antipsychotics (risperidone, quetiapine, sertindole, olanzapine, ziprasidone, aripiprazole) seem to represent an advance over first generation antipsychotic agents when treating psychotic symptoms in these patients due to a reduced risk of EPS. Clozapine should be used cautiously due to the risk of bone marrow suppression with agranulocytosis or lowered leukocyte levels. Lower starting doses, slower titration and close monitoring of side effects are recommended when treating HIV-infected patients with antipsychotic medication.
The WHO recommends that HIV-infected adults and adolescents should start ART therapy when there is WHO stage IV HIV disease, irrespective of the CD4 cell count, and in WHO stage III with consideration of using CD4 cell counts <350 cells/mm³ to assist decision-making. For those with WHO stage I or II HIV disease, a CD4 cell count of < 200 cells/mm³ or a total lymphocyte count of < 1200 cells/mm³ should trigger initiation of ART. The preferred regimen for initiating HAART under these circumstances are the combination of two nucleosides and one of the non-nucleoside drugs: stavudine or zidovudine (alternative: tenofovir or abacavir) plus lamivudine or emtricitabine, plus nevirapine or efavirenz. Triple NRTI approach can be considered as an alternative for first-line regimens in situations where NNRTI options provide additional complications. Computer programs for predicting response to combination antiretroviral therapy, given the viral genotype and further information, have lately been developed. Since the process of reverse transcription lacks a proofreading mechanism, mutations leading to drug resistance are generated frequently.

Efavirenz has well documented acute CNS side effects, and have been reported in up to 50% of the patients. Such side effects include dizziness, light-headedness, somnolence, insomnia, unusual dreams, amnesia, agitation, derealisation, depersonalization, hallucinations and euphoria, and typically begin after the first dose and usually resolve within 2-4 weeks of therapy or somewhat later. In certain patients, some symptoms may not completely resolve and persisting CNS side effects have been reported. Several other complications can occur during antiretroviral medication, for example metabolic and morphological changes such as lactic acidosis, lipatrophy, fat accumulation and dyslipidaemia. The immune reconstitution inflammatory syndrome (IRIS) is a spectrum of clinical signs and symptoms resulting from the restored ability to mount an inflammatory response associated with immune recovery. The incidence of IRIS is estimated to be 10% among all patients initiating antiretroviral therapy.

In the era of HAART, incidence rates of HAD have dropped by 15 to 50%, to as low as 10.5% or even lower. HAART can improve cognitive performance in some patients with HIV-associated cognitive impairment. It is not clear whether HAART protects the brain by restraining progress of the initial HIV infection in the CNS, or by controlling replication in the periphery and reducing the chance of later viral re-infection of the brain. Probably HAART acts in both ways.

However, HAART has failed to provide complete protection from HAD or reversal of the disease in many cases, and the frequency of the milder form of HIV-associated cognitive impairment, minor cognitive motor disorder (MCMMD) has not been reduced. Furthermore, the frequency of HIVE, the pathological manifestation of HAD and usually characterized only in post-mortem tissue, has remained constant. This suggests that HAART does not eliminate HIV infection in the CNS, although HAART improves the symptoms of HIV infection quite dramatically. HIV is believed to persist in sanctuary sites within the body even in optimally treated patients, and HAART is not a cure for HIV. The brain is believed to be one of the key sanctuary sites with limited immune surveillance and persistent HIV replication in microglial cells and perivascular macrophages. It has further been suggested, that the poor penetration of antiretroviral drugs across the blood-brain

**Antiretroviral therapy: the era of HAART**
The clinical manifestation of AIDS and HAD carries a poor prognosis for the affected patient. The introduction of highly active antiretroviral therapy (HAART) in 1986-1987 had a dramatic effect on HIV disease and increased the life expectancy for HIV-infected patients, and resulted in a 50% decline in AIDS death rate. Maternal-infant transmission rates have also decreased. The improved immune system protects against opportunistic infections and the incidence of such infections has fallen. Since the introduction of HAART, HIV infection in western countries has been transformed from an almost uniformly fatal condition to a chronic, manageable disease, even though some patients fail to achieve efficient viral suppression after receiving HAART due to drug resistance or lack of adherence.

HAART is now usually given as a combination of three or four different antiretroviral drugs from at least two different drug classes. There are over 20 different antiretrovirals approved, distributed in four different classes of antiretroviral drugs: nucleoside/nucleotide analogue reverse transcriptase inhibitors (NRTIs; e.g. abacavir, emtricitabine, lamivudine, stavudine, tenofovir, zidovudine), non-nucleoside analogue reverse transcriptase inhibitors (NNRTIs; e.g. efavirenz, nevirapine), protease inhibitors (e.g. lopinavir, ritonavir) and fusion inhibitors (enfuvirtide). The WHO has developed guidelines for the use of HAART in resource-limited settings.

**Substance use disorders**
There is a high prevalence of substance abuse among HIV infected individuals; studies have shown lifetime rates as high as 40-50%; in one study even as high as 74%. Researchers evaluating the relationship between substance abuse and HIV outcomes have primarily focused on injection drug users, but also alcohol, marijuana, amphetamines, cocaine and hallucinogens are frequently associated with HIV. In a recent study from South Africa, 35.8% of 536 patients attending HIV clinics reported alcohol abuse according to a screening test. Furthermore, drug and alcohol use disorders are frequently co-morbid also with other psychiatric diseases as depression, anxiety and severe mental illness. The combination of severe mental illness and substance use creates raised HIV risk, for example by increased sexual and needle-use risk-taking. HIV positive drug abusers are reported to have higher rates of both HAD and HIV compared to non-drug abusers. Up to 30% of injection drug users are HIV positive, and they seem to have more rapid neurologic progression. HIV related brain pathology thus seems to be worse in drug abusers compared to that seen in drug free patients. At present, there are no laboratory or clinical based methods to distinguish HIV mediated effects from drug abuse mediated effects on the brain. It has been shown that intravenous drug users with HAD do less well on HAART than non-drug users, and there might be greater difficulties in achieving compliance in drug users on antiretroviral therapy. At least 1 month should have elapsed since cessation of substance abuse before a diagnosis of HAD is made.

needed. The antiretroviral efavirenz may give rise to psychotic side effects (see below).

The clinical manifestation of AIDS and HAD carries a poor prognosis for the affected patient. The introduction of highly active antiretroviral therapy (HAART) in 1986-1987 had a dramatic effect on HIV disease and increased the life expectancy for HIV-infected patients, and resulted in a 50% decline in AIDS death rate. Maternal-infant transmission rates have also decreased. The improved immune system protects against opportunistic infections and the incidence of such infections has fallen. Since the introduction of HAART, HIV infection in western countries has been transformed from an almost uniformly fatal condition to a chronic, manageable disease, even though some patients fail to achieve efficient viral suppression after receiving HAART due to drug resistance or lack of adherence.

HAART is now usually given as a combination of three or four different antiretroviral drugs from at least two different drug classes. There are over 20 different antiretrovirals approved, distributed in four different classes of antiretroviral drugs: nucleoside/nucleotide analogue reverse transcriptase inhibitors (NRTIs; e.g. abacavir, emtricitabine, lamivudine, stavudine, tenofovir, zidovudine), non-nucleoside analogue reverse transcriptase inhibitors (NNRTIs; e.g. efavirenz, nevirapine), protease inhibitors (e.g. lopinavir, ritonavir) and fusion inhibitors (enfuvirtide). The WHO has developed guidelines for the use of HAART in resource-limited settings.

**Antiretroviral therapy: the era of HAART**
The clinical manifestation of AIDS and HAD carries a poor prognosis for the affected patient. The introduction of highly active antiretroviral therapy (HAART) in 1986-1987 had a dramatic effect on HIV disease and increased the life expectancy for HIV-infected patients, and resulted in a 50% decline in AIDS death rate. Maternal-infant transmission rates have also decreased. The improved immune system protects against opportunistic infections and the incidence of such infections has fallen. Since the introduction of HAART, HIV infection in western countries has been transformed from an almost uniformly fatal condition to a chronic, manageable disease, even though some patients fail to achieve efficient viral suppression after receiving HAART due to drug resistance or lack of adherence.

HAART is now usually given as a combination of three or four different antiretroviral drugs from at least two different drug classes. There are over 20 different antiretrovirals approved, distributed in four different classes of antiretroviral drugs: nucleoside/nucleotide analogue reverse transcriptase inhibitors (NRTIs; e.g. abacavir, emtricitabine, lamivudine, stavudine, tenofovir, zidovudine), non-nucleoside analogue reverse transcriptase inhibitors (NNRTIs; e.g. efavirenz, nevirapine), protease inhibitors (e.g. lopinavir, ritonavir) and fusion inhibitors (enfuvirtide). The WHO has developed guidelines for the use of HAART in resource-limited settings.

The WHO recommends that HIV-infected adults and adolescents should start ART therapy when there is WHO stage IV HIV disease, irrespective of the CD4 cell count, and in WHO stage III with consideration of using CD4 cell counts <350 cells/mm³ to assist decision-making. For those with WHO stage I or II HIV disease, a CD4 cell count of < 200 cells/mm³ or a total lymphocyte count of < 1200 cells/mm³ should trigger initiation of ART. The preferred regimen for initiating HAART under these circumstances are the combination of two nucleosides and one of the non-nucleoside drugs: stavudine or zidovudine (alternative: tenofovir or abacavir) plus lamivudine or emtricitabine, plus nevirapine or efavirenz. Triple NRTI approach can be considered as an alternative for first-line regimens in situations where NNRTI options provide additional complications. Computer programs for predicting response to combination antiretroviral therapy, given the viral genotype and further information, have lately been developed. Since the process of reverse transcription lacks a proofreading mechanism, mutations leading to drug resistance are generated frequently.

Efavirenz has well documented acute CNS side effects, and have been reported in up to 50% of the patients. Such side effects include dizziness, light-headedness, somnolence, insomnia, unusual dreams, amnesia, agitation, derealisation, depersonalization, hallucinations and euphoria, and typically begin after the first dose and usually resolve within 2-4 weeks of therapy or somewhat later. In certain patients, some symptoms may not completely resolve and persisting CNS side effects have been reported. Several other complications can occur during antiretroviral medication, for example metabolic and morphological changes such as lactic acidosis, lipatrophy, fat accumulation and dyslipidaemia. The immune reconstitution inflammatory syndrome (IRIS) is a spectrum of clinical signs and symptoms resulting from the restored ability to mount an inflammatory response associated with immune recovery. The incidence of IRIS is estimated to be 10% among all patients initiating antiretroviral therapy.

In the era of HAART, incidence rates of HAD have dropped by 15 to 50%, to as low as 10.5% or even lower. HAART can improve cognitive performance in some patients with HIV-associated cognitive impairment. It is not clear whether HAART protects the brain by restraining progress of the initial HIV infection in the CNS, or by controlling replication in the periphery and reducing the chance of later viral re-infection of the brain. Probably HAART acts in both ways.

However, HAART has failed to provide complete protection from HAD or reversal of the disease in many cases, and the frequency of the milder form of HIV-associated cognitive impairment, minor cognitive motor disorder (MCMMD) has not been reduced. Furthermore, the frequency of HIVE, the pathological manifestation of HAD and usually characterized only in post-mortem tissue, has remained constant. This suggests that HAART does not eliminate HIV infection in the CNS, although HAART improves the symptoms of HIV infection quite dramatically. HIV is believed to persist in sanctuary sites within the body even in optimally treated patients, and HAART is not a cure for HIV. The brain is believed to be one of the key sanctuary sites with limited immune surveillance and persistent HIV replication in microglial cells and perivascular macrophages. It has further been suggested, that the poor penetration of antiretroviral drugs across the blood-brain
HIV seems to have shifted from a subcortical pattern of neurocognitive impairment, reflecting a fluctuation of inflammatory processes during HAART treatment. The failure of this to respond to HAART. Such events may correspond to an irreversible stage of pathology with prominent neuronal loss. Determinants of treatment response are still unclear, but higher initial levels of CNS inflammation may correlate with reversible neurological deficits.

In the US, HAART has been associated with improvement in neurocognitive performance among HIV-positive individuals with a CD4 count of >200 cells/mm³ as well as CD4 count of ≥200 cells/mm³. This has to be further evaluated on sub-Saharan populations. However, WHO guidelines states that the presence of HIV clinical disease stage IV, for example HIV encephalopathy, should result in initiation of HAART, independent of CD4 count. The recognition of HIV-associated asymptomatic neurocognitive impairment (ANI) might also promote the initiation of antiretroviral therapy, independent of CD4 count or plasma HIV RNA levels, but this requires further study. Sacktor and colleagues studied the effect of HAART on cognitive impairment in a sub-Saharan cohort (23 HIV-positive individuals from Kampala, Uganda). There was a significant improvement in the Memorial Sloan Kettering HIV dementia stage and in tests of verbal memory, psychomotor speed and executive functioning after 6 months of HAART. The mean CD4 cell count improved from 71 cells/mm³ at baseline to 161 cells/mm³ at 3 months and 222 cells/mm³ at 6 months. The results from this study suggest that HAART should be provided for patients with HIV-associated cognitive impairment, also in resource-limited areas in sub-Saharan Africa.

HIV infection in older patients (aged >50 years) is becoming increasingly common as HIV-positive individuals live longer because of HAART treatment. The long-term consequences of chronic infection and extended exposure to HAART with respect to the brain are not known. Additive effects of for example drug abuse or co-infection with hepatitis C virus should also be considered. Several studies have identified an increased rate of HAD among older patients, but whether there is an additive or synergistic relationship between aging and HIV on neuropsychological performance is not clear. In addition, it is not known if HIV infection increases the risk for other age-related neurodegenerative disorders, perhaps by decreasing the age of onset and lowering the threshold for the clinical presentation. In another study by Sacktor and colleagues, the neuropsychological test performance between old and young HIV-positive cases were compared in patients with or without cognitive impairment, and in patients with dementia. Increased age was associated with lower performance in tests of executive functioning, memory and motor performance in patients with or without cognitive impairment, whereas older HIV-positive individuals with dementia may have greater decline in executive functioning compared to younger HIV-positive individuals with dementia. These differences could be a result of advance age itself or age-associated co morbidities such as coexisting neurodegenerative or cerebrovascular disease.

With HAART, dramatic reductions in plasma HIV levels can be seen within weeks, while immunological response occurs over a few months, and can be dramatic with normalization of CD4 counts. HAART also causes a reduction of HIV RNA levels in the CNS. Before HAART, traditional biomarkers such as CD4 level, plasma viral load, CSF viral load, and immune activation, e.g. monocyte chemoattractant protein-1 (MCP-1) and beta-microglobulin, were helpful in the diagnosis of HAD. However, during the era of HAART these biomarkers are less likely to be associated with dementia in treated HIV patients. For example, undetectable CSF HIV RNA do not indicate inactive HAD in HAART-treated patients. More sensitive assays are needed to reliable detect levels of HIV replication and active HIVE in the CNS. Bandaru and colleagues examined potential CSF biomarkers in patients with HIV, the majority on HAART, based on changes in cognitive status over a 1-year period. They found that increased levels of vitamin E and triglyceride CS2 predicted the onset or worsening of dementia, and that elevated levels of sphingomyelin was associated with inactive dementia. Elevated levels of ceramide and the accumulation of 4-hydroxynonenals were associated with active dementia. These findings indicate an initial up-regulation of the antioxidant defence early in the pathogenesis in HAD. The failure of this
defence mechanism leads to an accumulation of sphingomyelin. The breakdown of sphingomyelin to ceramide and the accumulation of reactive aldehydes are associated with a decline in cognitive function. Thus, a combination of markers of oxidative stress and neuronal injury might be useful in the future for following patients with HAD. However, these tests are currently available only in research laboratories.

Recent studies have shown that a marker of neuronal destruction, neurofilament light protein, is elevated in the CSF in patients with HAD. The possibility that HIV infection in the CNS might trigger some form of degenerative disease has initiated studies of markers typically associated with Alzheimer’s disease, and it has been shown that CSF amyloid beta42 and tau levels correlate with HAD.

Although the incidence of HAD has decreased since the introduction of HAART, the prevalence is increasing due to the rising number of infected subjects and increased life expectancy. The importance of HAD as an AIDS-defining factor has increased, and HIV neurocognitive disorders in the HAART era may occur even in those patients who do not have other evidence of active HIV disease. However, since the introduction of HAART, the course of HAD appears to be much more variable, and most HAART-treated patients with HAD remain neurologically stable, or may show some partial reversal of neurological impairment, for years after starting HAART. The milder form of HIV-associated cognitive impairment, MCDM, or MND, is now more common since the advent of HAART, and survival is considerably longer. An increased proportion of patients diagnosed with HAD now have a CD4 cell count >200 cells/mm³ and incomplete neuropsychological improvement following HAART has been described. Furthermore, the rate of HIV at autopsy has remained constant or might even be increasing. Thus, HAD continues to be an important cause of morbidity among HIV-positive patients also in the era of HAART, and significant cognitive impairment may persist within the limits of current treatment approaches. McArthur and colleagues suggest that HAD in the era of HAART may have three distinct subtypes: (1) a “subacute progressive” dementia in untreated patients similar to that seen in the pre–HAART era or in patients on HAART with high viral resistance; (2) a “chronic active” dementia in patients on HAART with poor adherence or with low viral resistance, and (3) a “chronic inactive” dementia in patients on HAART with good drug adherence and effective viral suppression. A fourth subtype, “reversible dementia”, has been described among patients with good drug adherence and effective viral suppression.

Neuroprotective therapies in the treatment of HIV-associated neurocognitive disorders has gained considerable interest, aiming to prevent brain injury or restore neurologic and neuropsychological function by mechanisms other than direct antiviral effects. Even if not as potent as HAART, they may diminish pathogenic events until antiretroviral therapy has reduced the viral burden, and they may provide mechanistic insight into CNS infection and brain injury. Adjunct benefit have been shown for some psychiatric medications. Two drug classes in particular, glycogen synthase kinase-3 beta (GSK-3β) inhibitors (valproic acid and lithium) and selective serotonin reuptake inhibitors (SSRIs) may be beneficial in this respect. Valproic acid (VPA), a histone deacetylase inhibitor, can induce activation of latent HIV within resting CD4 cells, possibly leading to an additional depletion of remaining provirus when used in combination with intensive ART therapy. VPA may also, in conformity with lithium, provide neuroprotection by inhibiting GSK-3β, thus protecting neural cells from lipid accumulation and apoptosis. Schiffito and colleagues conducted a phase II, double blind, placebo-controlled pilot study of VPA in HIV infected individuals. VPA was safe and well tolerated, and a small trend toward improvements in neuropsychological performance and brain metabolism was found in the VPA-treated subjects. Lithium has been demonstrated to prevent gp120-induced HIV neurodegeneration in vitro, an effect probably mediated by inhibition of GSK-3β. In a pilot study (single-arm, open-label, 12 weeks duration) low dose oral lithium significantly improved neuropsychological performance within eight cognitive impaired HIV patients already on ART.

SSRIs reduce the expression of CCR5, a chemokine receptor, limiting HIV entry into macrophages. In a recent study, HIV-positive individuals taking SSRIs were less likely to have detectable HIV RNA in CSF compared to non-users. This effect could not be attributed to better mood and adherence to ART.

Mementine, an uncompetitive antagonist of the N-methyl-D-aspartate (NMDA) receptor has been used in the treatment of neurodegenerative disorders as Alzheimer’s disease. It has recently been tested in a Phase II randomized, double-blind, placebo-controlled, multicenter trial on patients with mild to severe HAD and on stable antiretroviral therapy. Mementine was given for 16 weeks and was safe and tolerated by HIV-infected subjects, but no significant differences in cognitive performance was found, although magnetic resonance spectroscopy data suggested that mementine may ameliorate neuronal metabolism. The authors point out that longer studies are needed to assess the full potential of neuroprotective agents.

Some other agents have also been considered. The CNS stimulant modafinil tested on HIV-positive patients resulted in improved neuropsychological performance in an open-label, 4 weeks study. The psychostimulants methylphenidate and dextroamphetamine have both been studied in small placebo-controlled trials with some success in patients with HIV, and these observations justify further studies. The hypothesis that HAD is associated with oxidative stress and inflammation has led to preliminary studies evaluating the efficacy of selegiline (a dopamine agonist and antioxidant), minocycline (an antibiotic with anti-inflammatory effects), nimodipine (a calcium channel blocker) and TNF-α antagonists. There have been no controlled trials of cholinesterase inhibitors in treating dementia in HIV patients.

Research on the effect of HAART treatment of HIV positive patients with acute psychotic and/or manic symptoms is needed. In a recent pilot study, 42 HIV positive patients with psychiatric symptoms at the University of Cape Town, were given HAART ( stavudine and lamivudine with nevirapine or efavirenz). The dominating psychiatric manifestations were sub-acute delirium/organic affective psychosis. These patients...
had a high prevalence of multiple hallucinations, severe thought disorder, movement abnormalities, attention deficits, disorientation, subcortical cognitive impairment and symptom fluctuation. In addition, some patients had severe depression. Most of the patients responded very well to HAART. 16 patients were followed closely for 6 months, and showed a dramatic improvement in their psychiatric symptoms after 6-8 weeks on HAART. Furthermore, CD4 increased from 66.5 cells/mm³ at baseline to 256 cells/mm³ at 6 months. In addition to HAART, the patients were also given antipsychotic drug treatment, but more specific details of the treatment regime are not given in the report. Another study indicate that a CNS-penetrating antiretroviral (zidovudine) offer some protection against secondary mania.74

Conclusion

The use and distribution of HAART is steadily increasing outside the western world. The positive effects of HAART on HIV disease are well documented. HAART has also decreased the incidence of HAD, but does not give complete protection from this condition. To be able to follow and understand the clinical course of HAD, including possible subtypes, it will be important for future research to focus on the development of new biomarkers.31 There is also a need to determine the prevalence and development of HIV-associated neurocognitive impairment and other CNS complications of HIV, as well as confounders and co morbidities31, in patients who are receiving HAART.11,57 The therapeutic efficacy of HAART in the CNS is dependent at least in part upon its ability to achieve inhibitory drug concentrations. Therefore, finding new means of allowing drugs to cross the BBB and gain access to the CNS is critical for treatment of HAD and elimination of the brain as a potential viral reservoir.3,31 A better understanding of the pathogenesis of HAD is also needed to identify additional therapeutic strategies for the prevention and treatment of this neurodegenerative disease. The mechanism of HIV entry in the CNS, the pathological and neurotoxic events that occur in response to HIV CNS infection and the mechanism of viral evolution within the CNS should be further elucidated.3,9 Also, well designed clinical trials are required to gain a better understanding of how to treat HIV patients with psychotic and affective symptoms effectively and in safety.

References


