Factors Associated with Neuropsychological Impairment in HIV Infection

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1. Introduction

In today’s world, human immunodeficiency virus (HIV)/acquired immunodeficiency syndrome (AIDS) has poorly affected the vast majority of population and has been declared as one of the worst pandemic. It still remains the major public health burden across the world. The burden of HIV is increasingly progressing and due to lack of adequate knowledge among the population, their management is severely compromised. Advancement in the treatment of these cases are significant. New treatments are capable of reducing plasma viral load and in turn decreasing mortality (Carpenter et al., 1998). The betterment of patients with existing treatment regimen is quite evident, at least clinically and virologically. Here, important question is regarding life threatening impact of virus on various vital functions of the body. That includes central and peripheral nervous system (CNS) function. Such impact of virus becomes more evident, once their number increases to a numeric mark most often at the later stage of infection. However, it is reported that HIV enters the CNS within two weeks of initial infection (Krebs et al., 2000). Neuropsychological compromise are thought to occur secondary to HIV infection (Becker et al., 1997; Bornstein et al., 1993; Grant & A. Martin, 1994; Heaton et al., 1995; Hinkin et al., 1995; Maj et al., 1994a; Maj et al., 1994b; Martin et al., 1992; Stern, 1991; Van Gorp et al., 1989; von Giesen et al., 2001). When cases are presented, HIV-related neurocognitive disfunctioning may remain unnoticed at very early stage of infection. As the disease progress from asymptomatic to symptomatic and then to full blown AIDS, many neuropsychological impairments of clinical relevance appears in daily life of the patient, which may or may not have any functional significance. Especially, individual infected with HIV who is otherwise physically asymptomatic, these deficits may result in significant morbidity, in the sense that infected people may be unable to perform important tasks (e.g. bathing, cleaning, driving, work tasks), that influence their quality of life (QOL) and healthcare management directly. Such impairments may range from less severe unnoticeable defects to frank dementia syndromes...
that profoundly disrupt an infected individual’s functioning and activities of daily living. The neurocognitive deficits typically associated with HIV-1 infection include decrements in motor and information-processing speed, divided attention, memory retrieval processes, and executive functioning. Overall, the gross estimates of people infected HIV who experience neuropsychological impairment is range from 30% to 50% of total infected individuals (Grant & Martin, 1994; Foley et al., 2008).

Neuropsychological impairment can be classified into three subgroups (Grant et al., 1988). First group includes patients with subsyndromic neuropsychological impairment show deficits in at least two neuropsychological domains with no evidence of functional impairment. Second group is mild neurocognitive disorder or minor cognitive/ motor disorder and involve deficits in at least two domains (of ability) of neuropsychological test battery, sufficient affect patient daily functioning. Third group consist of most severe HIV-1 associated dementia which is characterized by moderate to severe cognitive and psychomotor slowing, impaired concentration and attention, memory disturbances and often motor incoordination and weakness. Many studies made attempt to found factors associated with neurocognitive impairment in HIV infection. However, factors related to neuropsychological impairment remain poorly understood. Herewith, we have reviewed the possible factors and divided them into three groups depending on their origin. Present chapter is an attempt to review the available literature on the possible factors and to discuss their mechanisms those are associated with neurocognitive impairment (Figure-1).

![Factors affecting HIV associated neuro-cognitive impairment](image-url)

Fig. 1. Factors affecting HIV associated neuro-cognitive impairment

HIV associated neuro-cognitive impairment /disorder are affected by three confounding factors; Biological factors, Psychological factors, Environmental and demographic factors. Initially, HIV does not easily get into the brain. But once it enters through blood-brain barrier and infects the resident microglial cells (a type of macrophage, which harbours the virus). Majority of the influence of biological factors are mediated by direct infection of HIV in brain. Symptoms of HIV associated neuro-cognitive impairments are misjudged with various other psychological problems like depression/ mood disorder, apathy and stress. Environmental and demographic factors like age, gender etc. also influence the HIV mediated neuro-cognitive impairment.
### 2. Biological factors

#### 2.1 Direct infection of HIV

Earlier, for long time, it was never thought of that HIV can also infect brain cells and that led to poor interpretation of many mental disorders associated with HIV-1 infection. Now, this biological stigma has been resolved. HIV-1 targets the cells of lymphoid and myeloid origin containing major HIV-1 receptors, CD4 and various chemokine receptors considered as HIV-1 co-receptors. These receptors help the attachment of the virus to the cell and the fusion of their membrane resulting in the entry of the virus into the cell (Zaitseva et al., 2003). Infected CD4+ T cells and monocytes, which circulate in the blood, are the potential source of CNS infection (Gonzalez-Scarano & Martin-Garcia, 2005). Many cells, such as T cells and monocytes are infected by HIV-1, these cells circulate in the blood and can cross the blood-brain barrier (BBB) and propagate the infection within the CNS (Figure-2) (Haase, 1986). A major part of HIV-1 associated neurological disfunctioning may be attributed to the entry of HIV-1 into the brain by crossing. The HIV-1 associated neuropathology is characterized by the infiltration of HIV-1 carrying macrophages into the CNS; the formation of microglial nodules; and multinucleated giant cells which result possibly from virus-induced fusion of microglia and/or macrophages in central white and deep gray matter; activation of astrocytes and their damage; neuronal loss particularly in hippocampus, basal ganglia and caudate nucleus. In addition, a variable degree of white matter pathology with evidence of broad range of myelin damage has been reported. HIV-1 has also been detected in the cerebral spinal fluid (CSF) (Gendelman et al., 1994). Imaging of brain using MRI, confirm that HIV infection is associated with progressive cortical atrophy within the gray and white matter, particularly in the later stage of the disease (Hall et al., 1996; Dal Pan et al., 1992; Stout et al., 1998; Aylward et al., 1993). These studies confirm that in the latter stage of infection, the anatomical atrophy in certain parts of brain occurs and many of them proven critical in the daily functioning of the individuals. These studies also report a correlation between the deterioration of cognitive function and the reduction in volume of certain brain structures.

#### 2.2 Immunodeficiency (opportunistic infections, CD4 & plasma viral load)

##### 2.2.1 Level of plasma HIV ribonucleic acid (RNA)

The levels of HIV-1 or its immune-dominant antigens/RNA (viral load) in plasma of infected individuals are the established marker for onset/progress of HIV related pathology. Infection of HIV-1 to brain cells is characterized by their presence in CSF. It is also proven that level of HIV RNA in CSF is elevated in subjects with cognitive impairments (Ellis et al., 1997; McArthur et al., 1997) and it also predicts future neuropsychological impairment (Ellis et al., 2002). The relationship between viral load and the mechanisms of cognitive decline requires further study. Clinically, however, a relationship between plasma HIV RNA levels and cognitive impairment would be more useful for diagnostic point of view. It is postulated that higher plasma HIV RNA levels may be associated with higher levels of circulating HIV-infected monocytes which may enter the blood-brain barrier and affect CNS functioning (Gartner, 2000). However, research on the direct relationship between plasma viral load and cognitive functioning remains equivocal. Various group failed to find a relationship between overall impairment and plasma HIV RNA (Ellis et al. 1997; Reger et al., 2005). However, contradictory reports are also available (Childs et al., 1999; Ellis et al., 2002). In a small neuropsychological battery, Stankoff et al. (1999) found a significant relationship
between motor functioning and plasma viral load. It seems that plasma viral load is associated with neuropsychological impairment but a lot is required to confirm its role as a predictive marker for neuropsychological impairment. Contradictory reports suggest that a comprehensive, in-depth and longitudinal approach with a multi-domain tool is needed is required to affirm its role as a marker.

2.2.2 Opportunistic infections
Opportunistic infections (OIs) of HIV patients significantly reduce their immunity to HIV and this majorly compromised their QOL. OIs generally occur in HIV infected patients with CD4 below 200, when their immune system is not sufficient enough to mount required immunity to prevent the OIs. Development of OIs further compromises their immunity, which in turn aggravates the expansion of HIV-1 and its spillage into CNS. It is also seen that presence of OIs in HIV infected patients are significantly associated with neuropsychological impairment in HIV infection. The determination of causes of neuropsychological impairment in HIV infection becomes more difficult because of various co-infections associated with it. More often heavy and chronic use of neuro-toxic drugs make it more complicated to study about exact cause. While challenging, these are questions that at one time in the HIV pandemic seemed difficult to imagine.

2.2.3 Treatment of disease
The effects of the HAART on neurocognitive functioning have not been assessed, although there have been some scanty studies showing neurocognitive effects of zidovudine (AZT) (Schmitt et al., 1988; Sidtis et al. 1993; Martin et al., 1999). Because drugs (HAART) can cross blood-brain barrier to achieve sufficient concentration to inhibit HIV replication in the CNS (Enting et al, 1998; Tashima, 1998; Limoges et al., 2000). It has also been shown that Zidovudine (AZT) monotherapy at current standard and higher doses can improve neuropsychological performance (Schmitt et al., 1988, Martin et al., 1999) and severity of dementia (Sidtis et al., 1993). However, the benefits of AZT monotherapy may be temporary because of the emergence of viral resistance (Gulevich et al., 1993). It was observed that individuals who are treated with HAART shortly after the first symptoms of dementia appear may show dramatic improvement.

2.2.4 CD4
Several studies reported that CD4 count is negatively related with neuropsychological impairment. Bornstein et al., (1991) found that patients with CD4 levels less than 200 had low scores on measures of motor speed, verbal memory acquisition, visual motor speed and mental tracking in comparison to patients with CD4 counts above 200. Osowiecki, et al. (2000) has reported a significant association of CD4 count with neurocognitive deficits. Perry, et al. (1989) found that among HIV positive patients, those scoring in the impaired range on neuropsychological measures had lower CD4 counts and lower CD4 /CD8 ratios than patients scoring in the normal range. It is also suggested that CD4 level correlates significantly with cognitive functioning. Several other studies have reported no relationship between neuropsychological performance and CD4 cell count (McArthur et al., 1989; Saykin et al., 1988; Martin et al., 1998). According to Honni and Bornstein (2002) there is very little evidence to suggest a significant association between CD4 count and the prediction of impairment.
Fig. 2. HIV crosses blood-brain barrier (BBB)

Infected macrophages (carrying HIV) crosses the blood brain barrier and enters into the brain and infects other cells. Infected cells as well as other resident brain cells produce copious amount of mediators (inflammatory factors, various neurotransmitter etc. In the presence of HIV, these factors irreversibly damage the astrocytes and neurons.

3. Psychological factors

3.1 Depression

Neuropsychological impairments are common in HIV infection with major depression although their nature remains partly unclear. Hypothalamic-pituitary-adrenal axis dysfunction is associated with neuropsychological dysfunction in major depressive disorder although evidence of direct causation is not definitive at present. Appearance of symptoms of depression has been significantly reported in patients with HIV-1 infection (Atkinson, et al., 1988; Perry, 1990; Perkins et al., 1994; Rabkin, 2000). However, the causative co-existence of major depression and neuropsychological impairment has not been reported. Additionally, symptoms of depression and neuropsychological impairment may occur together in many HIV-infected persons. Cysique et al., (2007) reported that neurocognitive impairment and major depression should be considered as two independent processes. Moreover, studies have demonstrated that neuropsychological abnormalities observed in HIV infection are distinct and cannot be attributed to depression (Marsh et al., 1994; Mapou et al., 1993; Perkins et al., 1994; Richardson et al., 1999; Bornstein et al., 1993, Claypoole et al., 1998; Goodwin et al., 1996; Goggin et al., 1997; Grant et al 1988; Moore et al., 1997). It has been shown that depressed patients with HIV-1 infection may exhibit deficits in learning and memory (Claypoole et al., 1998; Goggin et al., 1997; Kalechstein et al. 1998), but the contributions/involvement of depression to the impairment and severity of the
neuropsychological functioning appear to be minimal. However, more research is needed to confirm and widen these findings, and to expand the knowledge into clinical practice.

3.2 Apathy
Apathy refers to a cluster of symptoms reflecting lack of motivation manifested in motoric, emotional, and cognitive domains. To define here, motoric apathy is characterized by the tendency not to initiate a new motor activity unless externally prompted. Emotional apathy is defined as diminished intensity or persistence of emotion, or placidity, relative to the importance of some goal-directed thought or event. The third one, cognitive apathy is defined as indifference, a generalized loss of interest, decrease in goal-directed thought content diminished motivation associated with executive functions, and sometimes decreased verbal fluency. In context to the subject, Castellon et al., (1998) investigated the relationship between apathy, depression, and cognitive performance in HIV infection. They reported that apathy, but not depression, was found to be associated with working memory deficits among HIV subjects. Higher apathy scores and poorer working memory characterized the subjects with AIDS. Castellon et al., (1998) concluded that apathy is independent of depression and may indicate CNS involvement in HIV infection. In contrast, Rabkin et al., (2000) reported that apathy was consistently related to depression and unrelated to neuropsychological impairment.

3.3 Stress
Life stress has frequently been examined in the context of psychosocial factors in HIV-infected people because of the known adverse effects of stress on immunity of host (Patterson et al., 1995; Evans et al., 1997). In the context of cognitive function among HIV-infected people, the potential importance of stress has gained increased support from available literature. Moreover, these reports have demonstrated its adverse effects on brain structure and function. Several animal studies have reported atrophy in the CA3 region of the hippocampus in response to stress (Kim & Yoon, 1998). The chronic stress induced structural changes in brain is mediated by glucocorticoid hormones and neurotransmitters such as serotonin and the GABA-benzodiazepine system (McEwen, 2000). Furthermore, corticosterone a glucocorticoid appears to regulate levels of internal calcium (Ca2+) and thus may influence synaptic plasticity, aging, and cell death (Kim & Yoon, 1998). In context to patients, It has been demonstrated that atrophy in brain and abnormalities in its function are result of heightened levels of glucocorticoids and severe, traumatic stress (McEwen, 2000). In another study, it is observed that stressful life events were significantly related to cognitive performance only in the HIV-infected subjects (Pukay-Martin et al., 2003).

3.4 Drug addiction
It is well known that the intravenous administration of drugs with shared needles is an important cause for the spread of HIV burden. The persistent use of drugs through other routes (e.g., inhaling crack/cocaine) may also increase the risk for HIV infection. Drug abusers also practice prostitution to support their livelihood, habit and they may also engage in risky sexual behavior while under the influence of drugs or alcohol. But injection drug use with shared needle has also been seen as a main reason for the rapid progression of HIV among drug abusers (Bouwman et al., 1998), and has been found to diminish overall neuropsychological performance and reduce visuomotor processing, executive functioning,
motor speed and strength, and sensorimotor perception in persons in the very early stages of HIV infection (Claypoole et al., 1993). It was initially thought that frequent drug usage may impair the cognitive functioning among HIV-seropositive persons for (Wellman, 1992). It was also postulated that drug usage possibly induces CNS impairment independent of that caused by HIV infection. However, this argument does not appear to hold true, as most studies have found that neuropsychological dysfunction may not be contributed solely or partially by substance abuse, if we do not consider HIV seropositivity (Janssen et al., 1989). It was also suggested that the interaction of HIV infection and drug use might produce additive and synergistic cognitive deficits. Studies suggest that co-presence of HIV infection and drug use does not affect the cognitive functioning (Selnes et al., 1990; Selnes et al., 1992; Goodwin et al., 1996; Bornstein et al., 1993; Bono et al., 1996; Selnes et al., 1995). Cristiani et al., (2004) reported that the effect of marijuana use was greatest in subjects with symptomatic HIV infection. Further inspection suggested that this effect was due primarily to performance on memory tasks. Their findings suggest that although there is minimal impact of marijuana on uninfected individuals or those at early stages of HIV infection, there is a synergistic effect of HIV and marijuana use in patients with advanced HIV disease. Exception to the earlier studies, it was reported that HIV infection and methamphetamine dependence are each associated with neuropsychological deficits, and suggest that these factors in combination are associated with additive deleterious cognitive effects (Rippeth et al., 2004). It is also suggestive of that both HIV infection and methamphetamine dependence can be associated with brain dysfunction.

4. Environmental and demographic factors

4.1 Education

A certain level of education is required for the prevention and cure of HIV infection. It is now established that low education level is an independent risk factor for HIV-1 related cognitive impairment. Stern et al., (1996), Maj et al., (1994), Satz et al., (1993) reported that low educational level (6 years) increases the risk of HIV-1-related cognitive impairment in HIV-1-seropositive persons (asymptomatic and symptomatic). Our own unpublished data is in concordance with their findings. It is important to note that the association between low educational level and HIV-1-related cognitive impairment was independent of all other putative risk factors. In addition to this, another study also suggests that discrepancy in reading and education level is associated with worse neuropsychological performance (Ryan et al., 2005). There are two possible variables related to literacy, those are frequently used to predict the neuropsychological functioning; reading ability and years of education. In a country like India and particularly in its remote rural areas, it is difficult to predict the total year(s) of education. In such situations, it is better to rely on the reading ability of an individual. In other parts of world, previous research has also shown that reading ability is a stronger predictor of cognitive functioning than year(s) of education. Dotson et al., (2009) reported that reading ability predicts cognitive functioning than year(s) of education. They also suggest that influence of literacy and year(s) of education on cognitive functioning may vary among individuals belong to certain racial minority status and low SES group.

4.2 Age

The impact of age on some aspects of neuropsychological performance is well established. In addition, data indicate that age at the time of seroconversion is significantly associated
with HIV disease progression and survival time (Babiker et al., 2001). The possibility of an interaction between age and disease level on cognitive function has not been explored thoroughly. Both age and HIV status have been established as independent risk factors for the development of cognitive impairment, it is both reasonable and important to question whether the interaction of these two factors may constitute increased risk of impairment in HIV-infected individuals. While several studies have explored the impact of age on cognitive function in HIV infected subjects (Janssen et al., 1992; McArthur et al., 1993; Kim et al., 2001; Pereda et al., 2000) few have examined the interaction of age and disease status (Van Gorp et al., 1994; Hardy et al., 1999). Kissel et al., (2005) reported that Age and disease status had independent effects on cognitive function, but there were no significant interactions either on a summary measure of performance or on individual test scores.

4.3 Gender
Chiesi et al. and the AIDS in Europe Study Group (1996) reported that rates of cognitive impairment were higher for women. The hypothesis that female seropositive subjects are more vulnerable for neuropsychological impairment than seropositive men, is explained by Satz et al. (1993) that they usually have an extensive pre-morbid history, a history of substance abuse, a lower economic level, a greater psychiatric morbidity and a low educational level (Ickovics & Rodin, 1992; Melnick et al., 1994) variables that could increase the risk of neuropsychological impairment associated with HIV-1 (Stern et al., 1996). Failde-Garrido et al. (2008) reported different neuropsychological impairment pattern was detected between genders: while HIV+ men had greater impairment in visual memory, attention, psychomotor speed and abstract reasoning, HIV+ women had greater impairment on attention, psychomotor speed and verbal memory for texts. They further reported that there is no difference between the neuropsychological performance of seropositive male and female subjects. Pereda et al. (2000), Rabkin et al. (2000) found same result.

Fig. 3. Figure showing interrelationship among HIV infection, immunity of the host, neuropsychological functioning and quality of life (QOL)
HIV infection, immunity of the host, neuropsychological functioning and QOL are interlinked and affects each other. Various factors may influence neuropsychological functioning and immunity of the host. (-) and (+) sign shows negative association respectively. → and ↔ depicts uni and bidirectional interaction between two.

5. Conclusion

In conclusion, much can learn about factors associated with neuropsychological impairment, a leading cause of morbidity and mortality in persons with HIV infection. It was the intent of this manuscript not only to summarize what we know about factors associated with neuropsychological impairment in HIV infection but also to point out what we don't know and to highlight future directions that this area of research must now address. The adverse effect of HIV on neuropsychological functioning is well established. Manifestations of neurocognitive dysfunction range from subtle and mild cognitive changes to frank dementia syndromes. We have primarily focused on the factors associated with neuropsychological impairment which is adversely affect QOL and patients' ability demanding daily activities (Figure-3). There are very few studies focusing on factors associated with neuropsychological impairment in HIV infection. This review categorized factors into three broad areas that significantly associated with neuropsychological impairment. There are lacks of consensus among available literature regarding factors associated with neuropsychological impairment in HIV infection, especially psychological and environmental/demographic factors. Contradictory finding have been reported regarding association between neuropsychological impairment and antiretroviral therapy, drug abuse, depression, apathy, stress, age, gender and education. This review help the health care professionals in proper care of HIV seropositive people those are at risk to develop cognitive impairment. Present chapter also mentions the fact that tertiary prevention is urgently needed to improve quality of life of HIV infected people. Comprehensive study is needed to explore the above relationship and is clearly an important area of investigations.

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7. References


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The past few decades have seen the escalation of HIV-infections and the 'frantic' search for new drugs to treat the millions of people that live with HIV-AIDS. However because HIV-AIDS cannot be cured, but only controlled with drugs, and the Antiretroviral (ARV) treatment itself results in some undesirable conditions, it is important to generate wider awareness of the plight of people living with this condition. This book attempts to provide information of the initiatives that have been used, successfully or unsuccessfully, to both prevent and combat this 'pandemic' taking into consideration the social, economic, cultural and educational aspects that involve individuals, communities and the countries affected.

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