

Brain amyloid in virally suppressed HIV-associated neurocognitive disorder

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Abstract

Objective

To determine whether virally suppressed HIV neuropathogenesis, a chronic neuro-inflammatory state, promotes abnormal brain amyloid deposition.

Methods

A total of 10 men with virally suppressed HIV-associated neurocognitive disorder (HAND), aged 46–68 years, underwent ¹¹C-labeled Pittsburgh compound B PET. Data from the Australian Imaging, Biomarkers and Lifestyle (AIBL), including 39 cognitively normal individuals (aged 60–74 years), 7 individuals with mild cognitive impairment (MCI) (aged 64–71 years), and 11 individuals with Alzheimer disease (AD) (aged 55–74 years), were used as reference. Apart from more women, the AIBL cohort was demographically comparable with the HIV sample. Also, the AIBL PET data did not differ by sex. Cerebellum standardized uptake value ratio amyloid values within 22 regions of interest were estimated. In the HIV sample, apolipoprotein E (APOE) was available in 80%, CSF biomarkers in 60%, and 8–10 years of long-term health outcomes in 100%.

Results

HAND and the AIBL group with no cognitive deficits had similar amyloid deposition, which was lower than that in both the MCI and AD groups. At the individual level, one HAND case showed high amyloid deposition consistent with AD. This case also had a CSF-AD-like profile and an E4/E4 for APOE. Clinically, this case declined over 18 years with mild HAND symptoms first, followed by progressive memory decline 8–9 years after the study PET, then progression to severe dementia within 2–3 years, and lived a further 6 years. Another HAND case showed increased amyloid deposition restricted to the hippocampi. Two other HAND cases showed abnormally decreased amyloid in subcortical areas.

Conclusions

Relative to cognitively normal older controls, brain amyloid burden does not differ in virally suppressed HAND at the group level. However, individual analyses show that abnormally high and low amyloid burden occur.

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Glossary

AAL = Automated Anatomical Labeling; **A β** = amyloid beta; **AD** = Alzheimer disease; **AIBL** = Australian Imaging, Biomarkers and Lifestyle; **ANI** = asymptomatic neurocognitive impairment; **APOE** = apolipoprotein E; **^{11}C -PiB** = ^{11}C -labeled Pittsburgh compound B; **cART** = combination antiretroviral therapy; **HAD** = HIV-associated dementia; **HAND** = HIV-associated neurocognitive disorder; **MCI** = mild cognitive impairment; **MND** = mild neurocognitive disorder; **MSK** = Memorial Sloan Kettering; **NNTC** = National NeuroAIDS Tissue Consortium; **PLHIV** = people living with HIV infection; **PSP** = progressive supranuclear palsy; **PVE** = partial volume effect; **ROI** = region of interest; **SUVR** = standardized uptake value ratio.

The neuropathogenesis of HIV is commonly characterized by chronic neuroinflammation in people living with HIV infection (PLHIV) who are aging and virally suppressed on combination antiretroviral treatment (cART).¹ Both age and HIV-associated chronic inflammation may be associated with a higher risk of developing neurodegenerative disease such as Alzheimer disease (AD),¹ especially as PLHIV are entering age where dementia starts to rise in the general population.² AD and other forms of dementia and predementia syndromes are often characterized by abnormal deposition of cortical amyloid.^{3,4} This neuropathologic process may be used as a way to identify aging PLHIV at risk of dementia and may represent one of the endpoints of HIV-related chronic neuroinflammation. One of the most efficient detection methods for brain amyloid deposition in the living brain is the β -amyloid radioligand ^{11}C -labeled Pittsburgh compound B (^{11}C -PiB) positron emission tomography (PET).⁵ To date, there is no clear evidence that middle-aged PLHIV are more at risk of developing AD than their HIV-negative (HIV $-$) counterparts. In particular, it remains unclear whether HIV leads to AD or to an AD-like neurodegeneration for the following reasons.

Evidence from the literature remains limited and inconsistent. Achim et al.⁶ studied 40–65-year-old HIV+ participants using the molecular imaging probe 2-(1-{6-[(2-[F-18]fluoroethyl)(methyl) amino]-2-naphthyl} ethylidene)malononitrile (FDDNP)-PET and found increased binding of β -amyloid and tau protein aggregates in the posterior cingulate gyrus and the parietotemporal areas in three cases with mild neurocognitive disorder (MND). Furthermore, immunohistochemistry of the neocortex, subcortical white matter, and basal ganglia in HIV+ patients on cART found that 34 of 35 of these cases had intraneuronal deposition of β -amyloid. Tau and α -synuclein were also abundant with the distribution following axonal tracts. APOE genotyping, the major genetic AD risk factor, was not performed, and there were no concomitant neurocognitive data. Another study by Ances et al.⁷ that included 10 cognitively unimpaired HIV+ individuals (mean age 52 years, age range 46–58 years, 50% men) did not show an overall increased amyloid deposition using ^{11}C -PiB/PET compared with 20 HIV $-$ controls (age range 41–53 years, 80% men). This was despite 4 of 10 participants having low CSF amyloid beta (A β) 42 levels. The same research group led by Ances et al.⁸ also found no significant evidence of pathologic amyloid deposition with ^{11}C -PiB/PET when compared with HIV $-$ community age-comparable controls (88% men) in 5 HIV-associated neurocognitive disorder

(HAND) cases (82% men) and 11 cognitively unimpaired HIV+ patients (100% men) aged in their 40s. The HIV+ groups had a significantly lower amyloid deposition than 11 AD cases aged in their 70s (55% men). APOE genotyping was conducted in >90% of the HIV+ participants, and 45%–50% has at least one APOE 4 allele, but this was not associated with increased amyloid deposition. Analyses of CSF-AD markers were conducted in 77% of the HIV+ sample, with 5 HIV+ participants found to have low CSF-A β 42 levels (<500 pg/mL). CSF-A β 42 levels were not found to be associated with increased amyloid deposition. A larger study of 26 HIV+ adults aged 26–67 years old, reported as part of a review, found no difference in ^{11}C -PiB PET amyloid deposition and CSF-A β 42 levels between the HIV+ and the HIV $-$ patients aged 32–62 years old.⁹ This study did not provide APOE genotyping, and levels of cognitive performance were not reported. Finally, a case study of an HIV-infected (HIV+) 71-year-old man with an undetectable plasma viral load found a combined diagnosis of HAND and probable AD.¹⁰ This was based on a multidisciplinary examination including neuropsychological testing, brain imaging, CSF-AD markers, fluorodeoxyglucose PET, and florbetaben PET.

Overall, ^{11}C -PiB PET was the most commonly used method for detecting brain amyloid deposition. All studies conducted have focused on middle-aged PLHIV, mostly without HAND. All studies but one⁹ reported that most patients were on cART, although the degree of viral suppression was variable or not clearly reported. This represents a major caveat because detectable viral load has been found to be associated with increased A β -plaque in the hippocampus in a recent National NeuroAIDS Tissue Consortium (NNTC) analysis.¹¹ APOE genotyping was conducted in only one study.⁸ While APOE4 status has been associated with HAND in some studies^{12,13} this has not always been replicated,¹⁴ although this result may be influenced by cohorts that included PLHIV of relatively young age, who were not virally suppressed and clinically stable.¹⁴ Nevertheless, APOE4 status is a major risk factor for AD and other dementias¹⁴ and, thus, remains relevant in aging PLHIV cohorts that could be at higher risk of coincidental AD. Also, APOE4 is a strong modulator of ^{11}C -PiB PET data. In a cohort of 497 cognitively normal (NL) middle- and older aged participants with ^{11}C -PiB PET,¹⁵ a greater proportion of APOE4 carriers developed amyloid- β pathology, at an earlier age, and with faster amyloid- β accumulation. Furthermore, a recent and large neuropathologic study from the NNTC showed that APOE4 and age were strong independent

predictors of the deposition of 4G8-A β plaques in the frontal neocortex, evident in 29% of a selected sample.

¹¹C-PiB PET has high affinity and selectivity for fibrillar A β in plaques and extracellular amyloid deposits.⁵ It may thus lead to lower binding in PLHIV and patients with HAND for whom plaques have been found to be more often diffuse and intracellular.^{16–18} However, these neuropathologic findings were derived from cohorts that included relatively young patients, active AIDS, as well as many without full viral control and sometimes significant alcohol and drug use comorbidity.^{17–19} In addition, the detection of abnormal amyloid deposition in the oldest PLHIV with HAND using ¹¹C-PiB PET is important, as AD may be coincidental. On the other hand, the diffuse type of plaques may only reflect the early phase of the disease.

This study aimed to compare brain amyloid deposition quantified by ¹¹C-PiB PET in 10 formally diagnosed HAND cases who were part of the Australian HIV and Aging Cohort. HAND cases were screened for hepatitis C infection, non-HIV neurologic conditions, alcohol and drug use to minimize non-HIV-related neuroinflammation, and other forms of brain pathology.²⁰ These were compared with 39 NL cases, 7 cases with mild cognitive impairment (MCI), and 11 AD cases aged 55–74 years old, all from the Australian Imaging, Biomarkers and Lifestyle (AIBL) cohort.^{5,21} Because of the relatively small sample size and heterogeneous pattern of amyloid deposition in the HIV+ group, we also conducted a series of case analyses. Finally, we collected long-term health outcomes 8–10 years after the end of the PET data collection to determine dementia progression.

Methods

The University of New South Wales, St. Vincent's Hospital, and Austin Health Human Research Ethics Committees approved this research protocol (HREC/08/096). Informed written consent or the guardian consent when appropriate was obtained from all participants.

Participants

Ten HIV+ men formally diagnosed with HAND (table 1) were enrolled in a study to quantify brain amyloid deposition using the β -amyloid radioligand ¹¹C-PiB as part of the Australian HIV and Brain Aging Cohort Study between 2009 and 2011. The inclusion/exclusion criteria have been described in detail previously.²² All were on cART with undetectable plasma and CSF viral load (<50 copies/mL). Seven of 10 underwent a lumbar puncture and blood tests to determine CSF-AD markers and HIV markers, respectively, and 8 of 10 had APOE genotyping. All HAND cases underwent neuropsychological testing, and a high-resolution T1-weighted MRI scan was obtained for each person; these methods have been reported in detail elsewhere.^{13,22,23} Five cases had mild HAND, including 4 cases with asymptomatic neurocognitive impairment (ANI) and 1 case with MND. The 5 other cases

had HIV-associated dementia (HAD) along its clinical spectrum (Memorial Sloan Kettering [MSK] scale²⁴), including 1 case with mild HAD (MSK1), 2 with moderate HAD (MSK2), and 2 with severe HAD (MSK3). Note that on the MSK scale, MND is classified as MSK1.

AIBL data

A request was made to access the summed digital imaging and communications in medicine PET AIBL data on August 28, 2018, at aibl.csiro.au/research/support/ and was approved on August 30, 2018. These data were collected by the AIBL study group following a published protocol.²⁵ The AIBL study methodology has also been reported previously.^{5,21} AIBL subjects aged between 55 and 74 years with a PET scanning protocol closest to that of the HIV+ sample were retained (that is PET scan with a duration of 1,200 seconds: 57/103). Demographic and clinical characteristics of the retained AIBL cohort are presented in table 1. The AIBL sample has comparable demographics with the HIV sample except for more women, which is typical of an elderly cohort, because women have a higher survival rate and are more represented in AD cases²¹; HIV in Australia remains most prevalent in men who have sex with men.²² It is important that there was no significant differences between sexes in the AIBL PET data (*t* test *p* values >0.50). Although HIV infection was not formally diagnosed in those controls, the AIBL exclusion criteria were stringent and included a wide range of neurologic, psychiatric, and medical conditions in addition to laboratory test abnormalities.^{5,21} Evidence of immune compromise would therefore have been detected. Furthermore, the age and demographics of this cohort limits the risk of HIV infection because older age and late diagnosis are more common in non-Australian-born people.²⁶ See examples of AIBL PET data in figure e-1 and e-3, links.lww.com/NXI/A248, in supplemental material.

PET data acquisition in the HIV+ sample

All PET scanning took place at the Austin Health PET Centre in Melbourne (VIC, Australia). Patients were flown from Sydney and back, with their carer as appropriate. They were requested to fast for at least 6 hours before scanning. PET scans for the β -amyloid radioligand ¹¹C-PiB were acquired using a Philips Allegro PET camera with a resolution of 5.0 \times 5.0 \times 6.5 mm³ (*x y z*). A transmission scan was performed for attenuation correction. PET images were reconstructed using a 3D row action maximum likelihood algorithm with a voxel size of 2.0 \times 2.0 \times 2.0 mm³ (*x y z*). Summed images from the 40- to 70-minute time frame were used in each study. Each participant received 370-MBq ¹¹C-PiB IV over 1 minute. A 30-minute acquisition (6 \times 5-minute frames) in a 3D mode starting 40 minutes after injection of ¹¹C-PiB was performed (total PET scan duration of 1,800 ms).

PET processing and analyses in HIV and AIBL data

All imaging data sets were processed using the PETPVE12 toolbox²⁷ for SPM12 (v6906; Wellcome Trust Centre for Neuroimaging, London, UK; fil.ion.ucl.ac.uk/spm) in Matlab

Table 1 Demographic and clinical characteristics of the study samples

	HIV+	AIBL NL	AIBL MCI	AIBL AD
N	10	39	7	11
Age, y (range)	55.5 (46–68)	68 (60–74)	69 (64–71)	67 (55–74)
Male (%)	100	49	57	55
Education, y (range)	15 (8–19)	—	—	—
Urban dwelling MSM (%)	100	—	—	—
HIV duration, y (range)	13.8 (9.2–29.7)	—	—	—
Ethnicity (% white ESB)	100	—	—	—
Current cART duration, mo (range)	18.5 (6–120)	—	—	—
Nadir CD4 count, cells/mL (range)	203 (11–350)	—	—	—
Current CD4 count, cells/mL (range)	488 (210–1,008)	—	—	—
Current CD8 count, cells/mL (range)	1,074 (420–1914)	—	—	—
Predicted WAIS-III VIQ (range)	109.6 (84.6–121.1)	—	—	—
GDS (range)	1.7 (0.6–4.7)	—	—	—
Mini State Mental Examination (range)	—	30 (26–30)	27 (24–30)	22 (6–28)
Clinical Dementia Rating (range)	—	0 (0–0.5)	0.5	0.5 (0.5–3)
ANI, MND (count)	4, 1	—	—	—
HAD: MSK1, 2, 3 (count)	1, 2, 2	—	—	—
Historical AIDS (%)	70	—	—	—
Plasma HIV RNA undetectable (<50 copies/mL) (%)	100	—	—	—
CSF HIV RNA undetectable CSF (<50 copies/mL) (%)	100	—	—	—
Median CSF Aβ-42, pg/mL (range)	525.2 (142.5–663.3)	—	—	—
Median CSF t-tau, pg/mL (range)	207.5 (73.6–538.5)	—	—	—
Median CSF p-tau, pg/mL (range)	61.9 (22.6–115.9)	—	—	—
CSF-AD profile, % (count)	43 (3/7)	—	—	—
APOE (ϵ3/ϵ2)/ϵ4, % (count)	37.5 (3/8)	28.2 (11/39)	28.5 (2/7)	40 (4/10)
APOE ϵ4/ϵ4, % (count)	25 (2/8)	5.1 (2/39)	28.5 (2/7)	40 (4/10)

Abbreviations: AD = Alzheimer disease; AIBL = Australian Imaging, Biomarkers and Lifestyle; ANI = asymptomatic neurocognitive impairment; ESB = English-speaking background; GDS = global deficit score; HAD = HIV-associated dementia; MCI = mild cognitive impairment; MND = mild neurocognitive disorder; MSM = men who have sex with men; NL = cognitively normal; VIQ = WAIS-III Verbal IQ.

The AIBL is an Australian cohort that has recruited well-educated participants of white and English-Speaking background similar to that of the HIV cohort (see references 5 and 21 for further details).

Continuous data are presented as median and (range), unless otherwise specified. Percentage values provided for categorical data.

AIDS classification based on the Centers for Disease Control and Prevention (CDC) 1993 HIV disease staging system.

n = 7 HIV+ participants consented to lumbar puncture for CSF analysis.

n = 8 HIV+ participants were genotyped for APOE.

CSF-AD profiles are based on our publication¹³ and were (1) t-tau >350 and A β 1-42 <530 pg/mL; (2) p-tau >60 and A β 1-42 <530 pg/mL; and (3) t-tau >350 pg/mL and A β 1-42/p-tau <6.5.

Memorial Sloan Kettering (MSK) scale.²⁴ Note that the MSK1 = MND as per Frascati criteria.

r2012b (Mathworks Inc., Sherborn, MA), which has improved functions to handle amyloid PET partial volume effects (PVEs) compared with previous versions. First, all T1-weighted images were segmented and skull stripped using default parameters. Second, all PET images were coregistered to their associated T1 images using the toolbox's default parameters. PVE correction was performed using the Müller-

Gärtner method,²⁸ and the intensity of the PVE-corrected PET images was normalized to the average intensity of all cerebellar regions (cerebellum standardized uptake value ratio [SUVRc]) from the Automated Anatomical Labeling (AAL) atlas.²⁹ The inverse deformation parameters from the segmentation step were used to transform all AAL regions from the Montreal Neurological Institute to native space, separately

for all participants. Finally, SUVR_c values were extracted for each region of the AAL atlas in native space. To reduce the total number of regions of interest (ROIs), the 116 regions from the AAL atlas were arranged based on a previous atlas nomenclature²⁹ and averaged out to result in 22 final ROIs. Twenty-six regions related to the cerebellum and the vermis were used as the cerebellum reference. The remaining 90 regions were collapsed into the 22 ROIs. The algorithm used to collapse these regions was developed by M.S.K., neuroanatomist (see supplementary file table e-5, links.lww.com/NXI/A248).

Statistical analyses

Amyloid uptake and deposition distributions in the HIV+ sample

Two HIV cases had abnormal ¹¹C-PiB uptake (figure e-4, links.lww.com/NXI/A248) in which the PiB uptake was higher in the skull and some parts of the white matter compared with cortical regions. These were subsequently excluded based on the recommendation of the Austin PET team. As a result, 8 HIV cases were considered in the following analyses. The SUVR_c data were not normally distributed for any of the study groups. Therefore, all analyses were based on nonparametric statistics.

Group analyses

Study groups (HIV, NL, MCI, and AD) were compared using the Kruskal-Wallis test followed by the Steel-Dwass method to compare all pairs. There were no covariates entered in the analyses. Although the sex ratio did not significantly differ in the AIBL data ($p > 0.50$, and showed a slightly wider variance when women were included, table 2), the analyses were repeated excluding only the female participants in the AIBL NL group (N = 19) compared with the HIV sample, to account for a residual effect of sex. Owing to the limited sample size, this was not performed in the MCI (male = 4) and AD (male = 6) groups. These statistical analyses were conducted using the statistical package JMP 13 (SAS Institute Inc).

Case analyses

Case 2, on visual inspection, clearly exhibited high amyloid deposition across the entire brain. To quantify how significant the amyloid deposition was in this case, we ran case analysis statistics using the program Singlims_ES.exe^{30–32} against the AD, MCI, and NL groups (AIBL women included).

Case 1 had increased amyloid deposition in the hippocampus on visual inspection. Based on this, a single-case analysis was run for this ROI against the AD, MCI, and NL groups. Two other cases (cases 3 and 5) had a noticeable decrease in amyloid deposition across the basal ganglia. To quantify any significant differences, case statistics were run for these 2 cases for the cingulum, caudate, pallidum, putamen, and thalamus against the AD, MCI, and NL groups.

Long-term clinical outcomes in the HIV+ sample

We collected long-term clinical information from medical records located at St. Vincent's Hospital (Sydney, NSW,

Australia) and by contacting the participant's primary physicians. This allowed determination as to which patients had progressive dementia and other relevant comorbid conditions 8–10 years after the PET data collection.

Data availability

The HIV sample data are available on the UNSW Australia repository: ros.unsw.edu.au/ (search by author "Cysique" and title of the current study). The AIBL data are available upon request to the AIBL group at aibl.csiro.au.

Results

Group analyses

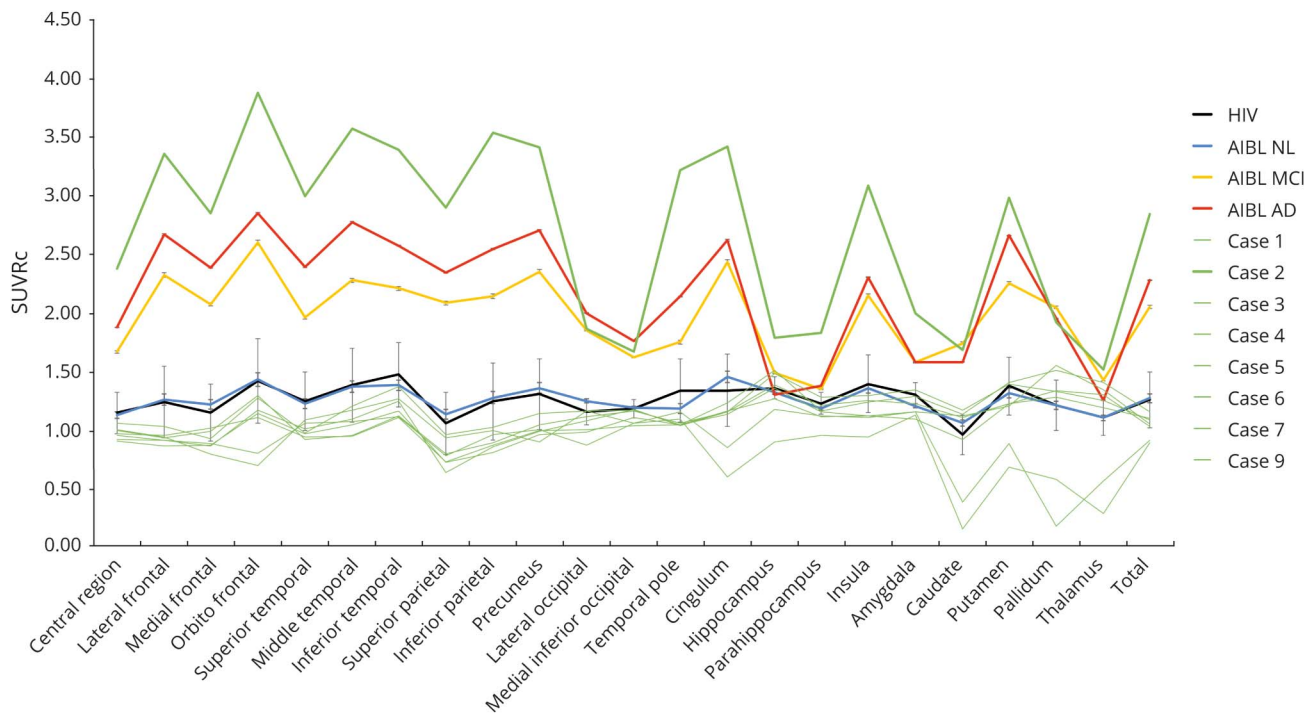
The HIV group did not differ statistically from the entire AIBL NL group on any comparisons. The HIV group showed a statistically lower amyloid deposition than the entire MCI group in the caudate and pallidum nuclei. Furthermore, the HIV group obtained a statistically lower amyloid deposition than the entire AD group in all regions, except for the hippocampi, amygdala, temporal pole, and thalamus. On all ROIs, the AD groups exhibited higher amyloid deposition than the NL group. The MCI group obtained higher amyloid deposition than the NL group on all regions, except for the thalami, hippocampi, occipital, parietal, and superior temporal regions (table 2 and figure 1). The analyses including only male participants produced similar results (table 2). Although because of a slightly more restricted variance in the AIBL NL male subgroup, we found a significantly lower amyloid deposition in the HIV sample in the medial frontal and superior parietal cortex.

Case analyses

On individual inspection, case 2 (aged 66 years) had high amyloid deposition consistent with AD (whole-brain SUVR_c = 2.83) (figures 1 and 2). On quantitative single-case analysis, case 2 showed a significantly increased deposition compared with the NL group, but no difference when compared with the MCI and AD cases. When including only men in the analyses, significant differences were again found between case 2 and the NL and AD groups, but not compared with the MCI group. These results are compiled in table 3, and individual ¹¹C-PiB PET images are provided (figure 2) and show that case 2 had the highest amyloid deposition compared with all the other groups. As described in table 3, case 2 was APOE ε4 homozygous with a CSF profile, based on standard cutoff values, consistent with AD. A detailed clinical history and neuropsychological profile for case 2 is presented in table 2. It shows a mix of mild and moderate impairment in cognitive domains affected by HIV-related brain injury (attention/working memory and speed of information processing), but not for motor functions. The memory profile was consistent with probable AD with moderate to severe impairment.

Case 1 (aged 62 years) had elevated amyloid deposition in both hippocampi (table e-1, links.lww.com/NXI/A248). Single-case analysis showed that there was no significant

Figure 1 PiB SUVR_c in the 22 ROIs in the study samples and individually in the HAND cases



The HIV+ group data are represented by the black line, the AIBL NL group data are represented by the blue line, the AIBL MCI group is represented by the yellow line, and the AIBL AD group is represented by the red line. Mean and SE were provided for all 22 brain regions and total brain SUVR_c. Individual HIV data were also represented by the green line, with the case who had amyloid deposition in line with AD in a bolded green line. AD = Alzheimer disease; AIBL = Australian Imaging, Biomarkers and Lifestyle; HAND = HIV-associated neurocognitive disorder; MCI = mild cognitive impairment; NL = cognitively normal; ROI = region of interest; SUVR_c = standardized uptake value ratio with reference to the cerebellum.

increase in deposition compared with the MCI and AD groups, but there was a trend compared with the NL group ($p = 0.06$; table 2 and figure 2). This individual was also APOE $\epsilon 4$ homozygous; however, CSF biomarkers were not consistent with an AD profile (CSF A β -42 = 168.21 pg/mL; t-tau = 132.18 pg/mL; and p-tau = 36.23 pg/mL). Neuropsychological deficits included mildly impaired speed of information processing and working memory and moderately impaired motor functions with preserved verbal fluency and memory.

Case analysis statistics, as well as demographic, neuropsychological, HIV, CSF, and APOE data for case 3 (aged 52 years) and case 5 (aged 54 years) are presented in table e-2, links.lww.com/NXI/A248, and e-3, links.lww.com/NXI/A248, respectively, and in figure 2. There was a significant decrease in amyloid deposition for both cases in all areas tested (range $p < 0.04$ – $p < 0.0001$), except for the cingulum and putamen when compared with the NL group ($p = 0.15$ – 0.07). Furthermore, case 5 was identified as having an AD-like CSF profile. Both were cases with severe HAD.

Long-term follow-up

The 2 patients who originally had clinically severe HAD (MSK3) and 1 who had clinically moderate HAD (MSK2)

and had normal amyloid died because of other comorbidities (see table e-4, links.lww.com/NXI/A248). Case 2 who had clinically moderate HAD + AD (MSK2) and significantly elevated amyloid also died after progression to severe dementia. Case 1 who had ANI, with elevated amyloid in the hippocampi, developed a progressive supranuclear palsy (PSP)-like illness, which has remained static. Case 1 also showed microinfarcts and hyperintensities of the white matter consistent with vascular injury on MRI and evidence of cognitive decline reaching an MND (MSK1) diagnosis within 3 years after study. Other cases, which had normal amyloid, included 1 ANI case (case 9), who did not progress clinically according to their physicians and are currently living independently. Case 6, who had MND (MSK1) at entry, has been cognitively stable despite a series of cancers, which have been successfully treated. Case 7, who had clinically mild HAD (MSK1) at entry, has been mostly stable except for progressing peripheral neuropathy. Of the 10 cases, 2 cases (cases 8 and 10) showed abnormal PiB PET uptake (figure e-4, links.lww.com/NXI/A248). Case 8 had ANI at entry that progressed clinically displaying worsening mood changes and behavioral problems with subsequent cognitive decline consistent with MND (MSK1). Case 10 also progressed clinically by reporting increasing cognitive difficulties in line with MND (MSK1). These 2 cases have not reached the HAD stage and still live independently.

Table 2 Comparisons of amyloid deposition on the study groups

ROI	HIV+	AIBL NL	AIBL NL male	AIBL MCI	AIBL AD	p1	p2	p < 0.05 on Steel-Dwass
Total	1.04 (0.12)	1.08 (0.22)	1.08 (0.13)	2.17 (0.70)	2.40 (0.43)	<0.0001	0.37	d, e, f
Central region	0.95 (0.07)	1.01 (0.16)	1.00 (0.14)	1.78 (0.84)	1.91 (0.37)	<0.0001	0.22	d, f
Lateral frontal	0.92 (0.04)	0.99 (0.30)	1.00 (0.17)	2.44 (0.86)	2.73 (0.42)	<0.0001	0.11	d, e, f
Medial frontal	0.87 (0.09)	1.02 (0.29)	1.03 (0.31)	2.25 (0.79)	2.49 (0.35)	<0.0001	<0.05	d, e, f
Orbitofrontal	1.13 (0.27)	1.14 (0.21)	1.13 (0.20)	2.71 (0.53)	2.92 (0.53)	<0.0001	0.87	d, e, f
Superior temporal	0.96 (0.08)	1.02 (0.17)	1.03 (0.16)	2.19 (0.91)	2.36 (0.68)	<0.0001	0.46	d, f
Middle temporal	1.05 (0.13)	1.11 (1.16)	1.10 (0.15)	2.64 (1.04)	2.84 (0.72)	<0.0001	0.20	d, e, f
Inferior temporal	1.15 (0.14)	1.18 (0.21)	1.19 (0.19)	2.36 (0.82)	2.51 (0.64)	<0.0001	0.87	d, e, f
Superior parietal	0.77 (0.14)	0.91 (0.30)	0.99 (0.23)	2.36 (1.65)	2.32 (0.52)	0.0001	0.04	d, f
Inferior parietal	0.87 (0.11)	1.02 (0.26)	1.04 (0.12)	2.62 (1.34)	2.62 (0.54)	<0.0001	0.02	d, f
Precuneus	0.98 (0.05)	1.10 (0.28)	1.12 (0.20)	2.66 (1.16)	2.69 (0.49)	<0.0001	0.10	d, e, f
Lateral occipital	1.07 (0.15)	1.12 (0.28)	1.07 (0.19)	1.95 (0.86)	1.88 (0.60)	<0.0001	0.15	d, f
Medial inferior occipital	1.09 (0.11)	1.11 (0.16)	1.11 (0.21)	1.55 (0.63)	1.65 (0.39)	<0.0001	1.0	d, f
Temporal pole	1.03 (0.03)	1.05 (0.14)	1.06 (0.18)	1.69 (0.45)	2.24 (0.71)	<0.0001	0.79	e, f
Cingulum	1.14 (0.17)	1.22 (0.21)	1.23 (0.19)	2.42 (0.78)	2.71 (0.27)	<0.0001	0.15	d, e, f
Hippocampi	1.34 (0.21)	1.32 (0.14)	1.32 (0.16)	1.42 (0.24)	1.31 (0.25)	0.07	0.63	f
Parahippocampi	1.14 (0.07)	1.14 (0.10)	1.13 (0.09)	1.25 (0.23)	1.36 (0.20)	0.0008	0.67	e, f
Insula	1.10 (0.13)	1.18 (0.26)	1.16 (0.16)	2.14 (0.38)	2.28 (0.62)	<0.0001	0.91	d, e, f
Amygdala	1.14 (0.11)	1.15 (0.09)	1.15 (0.08)	1.50 (0.25)	1.65 (0.40)	<0.0001	0.67	e, f
Caudate	1.10 (0.48)	0.99 (0.30)	0.95 (0.33)	1.66 (0.53)	1.37 (0.59)	<0.0001	0.59	b, d, e, f
Putamen	1.21 (0.26)	1.19 (0.25)	1.15 (0.15)	2.33 (0.40)	2.56 (0.46)	<0.0001	0.42	d, e, f
Pallidum	1.31 (0.49)	1.15 (0.33)	1.13 (0.28)	2.02 (0.25)	2.07 (0.73)	<0.0001	0.25	b, d, e, f
Thalamus	1.24 (0.44)	1.11 (0.23)	1.11 (0.25)	1.35 (0.34)	1.22 (0.37)	0.03	0.25	f

Abbreviations: AD = Alzheimer disease; AIBL = Australian Imaging, Biomarkers and Lifestyle; MCI = mild cognitive impairment; NL = cognitively normal; ROI = region of interest; SUVRc = standardized uptake value ratio with reference to the cerebellum. SUVRc values presented as median (IQR).

p1 is the overall p value extracted from the Kruskal-Wallis tests.

p2 is the Wilcoxon p value for the comparison between the HIV sample and male NL from AIBL.

p < 0.05 on the all pairs, Steel-Dwass method multiple comparisons: (a) NL vs HIV; (b) MCI vs HIV; (c) MCI vs AD; (d) HIV vs AD; (e) NL vs MCI; and (f) NL vs AD; ns = not significant at p < 0.05 on all comparisons.

Discussion

Consistent with previous studies,⁷⁻⁹ there was no statistical difference in amyloid deposition between the HIV group and NL cases (combining men and women) from the AIBL. However, the NL cases, as part of the AIBL study, were older than those in previous referenced studies. When repeating the analyses with only NL male participants from the AIBL, we found abnormally low amyloid deposition in the HIV group in the medial frontal and superior parietal cortex. This indicates that HIV may be associated with abnormally low amyloid deposition when compared with an older NL cohort. Our study has a small HIV sample size, yet it was a data-rich

sample composed of well-characterized HIV+ cases aged 55.5 (46-68) years old, with formally diagnosed HAND, long-term HIV disease clinical stability, viral suppression, and low neurologic and psychiatric confounds. Critically, the sample had long-term follow-up of the participants over years (table e-4, links.lww.com/NXI/A248). This is unique compared with previous HIV PET studies. The small HIV sample size was amenable to a case analysis strategy, which showed abnormal increases of amyloid deposition in 2 HAND cases.

Case 2 had an overall deposition of amyloid consistent with AD supported on single-case analysis. The typical AD-like CSF profile and progressive cognitive impairment with a mixed

Table 3 Case 2 with a probable AD and MND neuropsychological profile concomitant to the PET examination and clinical history

Demographic and clinical data	
Age, y	66
Education, y	18
HIV duration, y	24.6
Current cART duration, mo	21
Nadir CD4 count, cells/mL	18
Current CD4 count, cells/mL	210
Current CD8 count, cells/mL	882
Predicted WAIS-III VIQ	121.1
GDS	1.25
Historical AIDS	Yes
CSF A β -42	142.5
CSF t-tau	328
CSF p-tau	71.7
CSF-AD-like profile	Yes
APOE	ϵ 4/ ϵ 4
Total SUVRc vs NL	<0.0001 (3.60) (2.77–4.52) ^a
Total SUVRc vs MCI	0.11 (1.31) (0.30–2.45)
Total SUVRc vs AD	0.07 (1.60) (0.72–2.60)
Total SUVRc vs NL (males only)	0.002 (3.30) (2.12–4.57) ^a
Total SUVRc vs MCI (males only)	0.10 (1.58) (0.08–3.40)
Total SUVRc vs AD (males only)	0.04 (2.18) (0.71–3.96) ^a
Neuropsychological data	
Cognitive domains	Degree of cognitive impairment
Attention/working	Mild impairment
Speed of information processing	Mild impairment
Mental flexibility	Mild to moderate impairment
Verbal learning	Mild to moderate impairment
Verbal active retrieval	Moderate to severe impairment
Verbal memory retention	Moderate to severe impairment
Verbal fluency	Within normal range
Motor	Within normal range

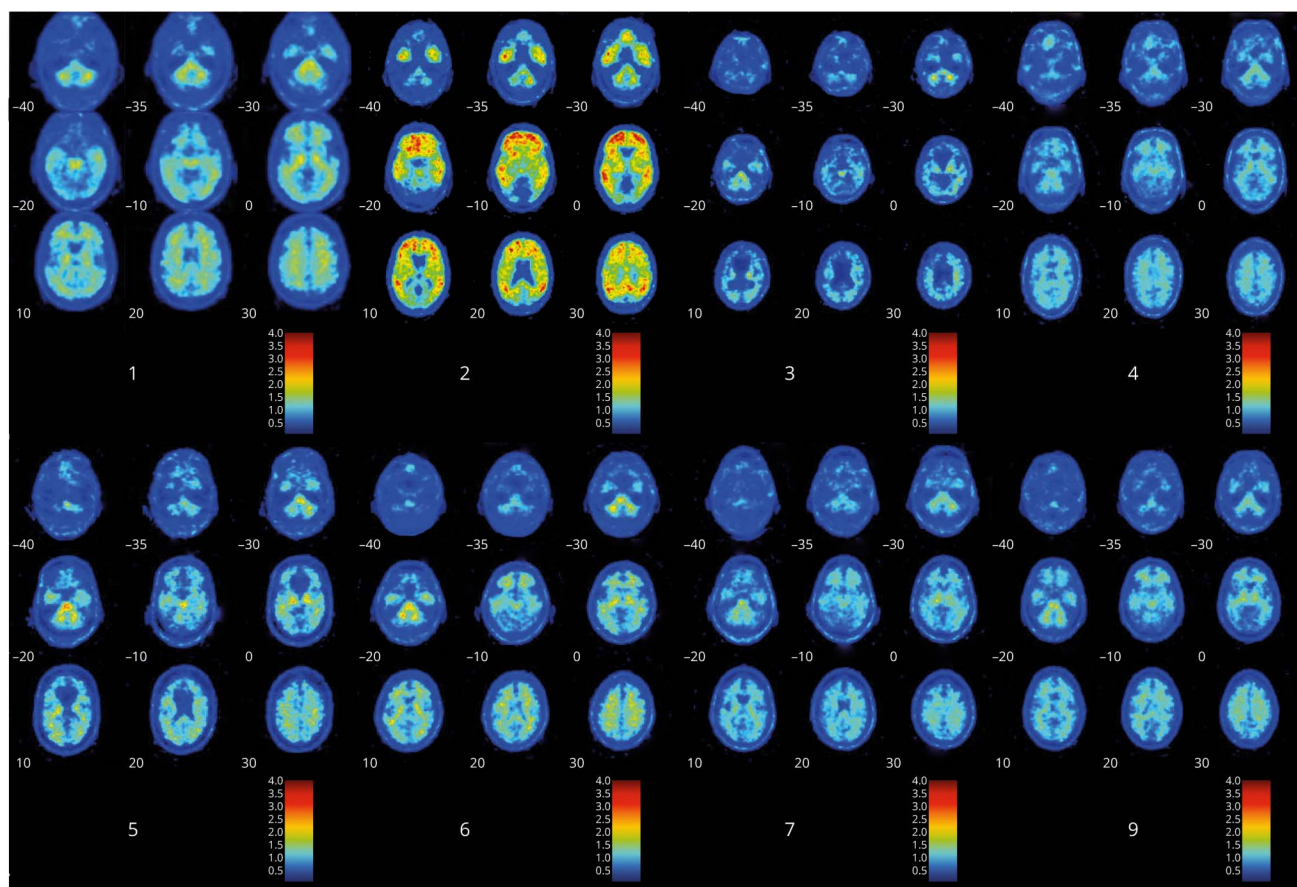
Abbreviations: AD = Alzheimer disease; GDS = global deficit score; HAND = HIV-associated neurocognitive disorder; MCI = mild cognitive impairment; MND = mild neurocognitive disorder; NL = cognitively normal; SUVRc = cerebellum standardized uptake value ratio; VIQ = WAIS-III Verbal IQ. Case statistics reported as 1-tailed *p* value (t value) (confidence interval).

The reported neuropsychological data were normed against local normative data.²²

This case has been in the clinical care of Sydney St. Vincent's Hospital HIV clinics for more than 25 years. His cognitive decline was documented across 18 years through standard neurologic (CSF, blood, and MRI/MRS examinations) and neuropsychological examinations at the neurology department. In 2001, this case was diagnosed with mild HAND in which symptoms remained stable for the next 10 years (mild learning and mild mental flexibility impairment). His premorbid abilities were in the superior range. His progressive memory decline started 8–9 years after his initial HAND diagnosis (~1 year ahead of the PET) as shown in the table above. At this time, he showed no apraxia, no agnosia, and no reasoning deficits (data from a standard clinical neuropsychological assessment performed during this period), in addition to no language and no motor deficits for the research examination as shown in the table. It is not until the PET examination that AD was suspected. Within 2–3 years, he progressed to severe dementia; this started with a decline in attention/working memory, speed processing, and further learning/memory decline. He lived for 6 years at this stage.

^a *p* < 0.05.

Figure 2 Raw PiB PET data in the 8 HIV-associated neurocognitive disorder cases



Raw PiB PET data are presented in the 8 HIV cases. Case 2 is the case with high amyloid deposition. Case 1 is the case with a trend for higher hippocampal binding compared with the Australian Imaging, Biomarkers and Lifestyle NL group. Cases 3 and 5 were the cases with the lowest amyloid deposition across all groups. Note that case example for the NL, MCI, and Alzheimer disease groups is presented in supplemental material in figures e-1, e-2, and e-3, respectively, links.lww.com/NXI/A248. MCI = mild cognitive impairment; NL = cognitively normal.

picture of typical AD and HAND deficits supported a probable dominant AD diagnosis. It is important that case 2 was APOE E4 homozygous and was the second oldest participant in the sample (66 years of age). This is consistent with potential coexistence of HAND and AD in an advanced HIV+ patient, as previously described.¹⁰ From an AD prevalence perspective, it is logical that some HIV+ patients will develop AD based on genetic and other predisposing factors. It is, however, not clear whether the start of neurodegeneration may be earlier because of HIV. It is also uncertain whether amyloid deposition may have a faster deposition rate,³³ although this is certainly possible given that this case had the highest amount of amyloid deposition across all our comparisons. Also, we noted that the cognitive profile was unusual compared with HAND (sparing of motor functions and long-term stability of psychomotor speed) and AD profiles (restricted memory deficits with no agnosia, apraxia, or language deficits except toward the end stage).³⁴ From a clinical perspective, it has been argued that HAND and AD can be distinguished to some degree.² However, an extensive test battery with the inclusion of agnosia, apraxia, and language would be needed in addition to the usual cognitive domains assessed in HAND to better address this

question. It has been noted that ¹¹C-PiB PET, which has high affinity and selectivity for fibrillar A β in plaques and extracellular amyloid deposits,⁵ may lead to lower binding in PLHIV and patients with HAND for whom plaques have been found to be more often diffuse and intracellular.^{16–18} In this study, case 2 shows that such an interpretation may not be correct in aging PLHIV with a possible coincidental AD pathology and profile.

A role for HIV in the dysregulation of brain amyloid is biologically plausible. Hypotheses include recent research into the antimicrobial activity of amyloid^{35,36} in which increased amyloid may reflect a long-term response to the presence of HIV infection in the brain (potentially in brain cellular reservoirs). Supporting this, Bourgade et al.^{37,38} suggest that β -amyloid can inhibit herpes simplex virus 1 and is associated with overproduction of amyloid more generally in infection-related neurologic diseases.

Although case 1 showed no statistical overall increase in amyloid deposition, a trend toward abnormal binding in the hippocampus was observed. The CSF biomarker profile was also not consistent with an AD-like profile. Recent literature on AD

suggests that, even subthreshold increases in amyloid levels can result in memory deficits, and in some cases it is the rate of amyloid accumulation, rather than the overall amyloid level, that more accurately predicts worsening memory.^{39,40} However, case 1 had a PSP-like illness, the pathologic underpinning of which in the context of HIV is unknown at present. Nonetheless, it is interesting that he was homozygous for APOE E4. There is evidence that progression into a PSP-like illness is associated with a homozygous APOE E4 genotype.⁴¹

In this study, 2 of the 3 cases (cases 3 and 5) with severe HAD (pre-cART) who survived on cART showed low amyloid deposition. These cases were the most vulnerable and died within 8–10 years of the study's long-term outcomes collection. All were younger than 65 years with long HIV duration (>25 years). Cases 3 and 5 exhibited major subcortical atrophy, consistent with severe HIV-related brain injury.^{42–45} This level of subcortical atrophy potentially biased the measurement of amyloid in the basal ganglia. Nevertheless, there is at least 1 plausible mechanism for the abnormally low level of amyloid. Microglia and astrocytes primed by HIV may play important roles in engulfing and purging A β plaques from the brain.⁴⁶ In severe HAD with a disrupted blood-brain barrier, amyloid clearance may be enhanced in some individuals. Also, in treated PLHIV, amyloid clearance may occur through HIV-1 Tat priming and activation of microglia.⁴⁷ Case 5 had a CSF biomarker profile that supported AD (T-tau >350 Ab42/P-tau <6.5), but the PET scan did not confirm increased amyloid deposition suggesting caution in the clinical use of CSF for AD diagnosis in patients with HIV, especially those with HAND. Finally, we cannot exclude that in these younger cases, the plaques may have been diffuse rather than fibrillar, and this may have contributed to a lower ¹¹C-PiB PET signal.^{16–18}

Although our study does not provide statistical evidence of an overall increase in amyloid in PLHIV with HAND, it highlights the importance of interindividual variations in patients with chronic HIV and HAND and the need to focus the research on favorable and unfavorable phenotypic and/or biotypic clusters.⁴⁸ In this small sample, we were able to detect cases with high, focused (albeit trending) and very low brain amyloid. Such complex phenotypes if already apparent in a pilot study suggest a need to focus on long-term individual phenotypic analyses. Future longitudinal studies monitoring amyloid deposition as PLHIV age are crucial in understanding the relationship between abnormal amyloid levels and HAND as well as other comorbidities. It is noteworthy that all the HIV+ patients in this study also had a large number of medical comorbidities (See table e-4). How exactly such comorbidities played a role in their amyloid pattern and then clinical progression or lack thereof remains to be fully elucidated. Although we used a PET protocol that retained the individual neuroanatomy, it would be important that larger studies use a multimodal imaging strategy because abnormal brain aging in HIV can be detected by using volumetric MRI.⁴⁹

Our study had some limitations. The controls used in this study were obtained from AIBL and included a mix of male and female participants differing from the HIV+ sample who were all men. The sex difference had a minimal effect on the cases with AD + HAD, when specifically investigated. We also had missing data for the APOE genotyping and CSF profiles and recognize this, just as for previous studies have, as a limitation. Although 50% of the HAND cases had HAD as per Frascati criteria at study entry, it is important to note that the MSK clinical staging was well represented as well as those with mild HAND as per Frascati criteria. Finally, our pilot sample size and case series analyses yield indicative rather than definitive findings. It remains unclear why 2 HAND cases had no cortical amyloid uptake. Despite a thorough search, we could not find reports of similar issues in the HIV or non-HIV literature. There was no PET scan technical issue reported by the Austin Health PET Centre examiners.

In conclusion, relative to NL older controls, brain amyloid burden does not differ in virally suppressed HAND at the group level. However, individual analyses show that abnormally high and low amyloid burden occur. These results indicate that focus on individual phenotypes in longitudinal analyses in the HIV-aging cohort is needed.

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Mustafa S. Kaseem, PhD	NeuRA	Manuscript writing
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