Brain PET in the Diagnosis of Alzheimer’s Disease

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Objectives: The aim of this article was to review the current role of brain PET in the diagnosis of Alzheimer dementia. The characteristic patterns of glucose metabolism on brain FDG-PET can help in differentiating Alzheimer’s disease from other causes of dementia such as frontotemporal dementia and dementia of Lewy body. Amyloid brain PET may exclude significant amyloid deposition and thus Alzheimer’s disease in appropriate clinical setting.

Conclusions: FDG-PET and amyloid PET imaging are valuable in the assessment of patients with Alzheimer’s disease.

Key Words: FDG brain PET, amyloid brain PET, dementia, Alzheimer’s disease

(Clin Nucl Med 2014;39: e413–e426)

LEARNING OBJECTIVES

After completing this journal-based CME activity, participants will be able to:

1. Identify patterns of FDG spatial distribution in brain PET of patients with MCI, AD, frontotemporal dementia, and dementia of Lewy Body.
2. Identify the value and learn the appropriateness criteria for amyloid brain PET.

Dementia is a collective group of neurodegenerative disorders characterized by memory impairment and cognitive decline, leading to difficulty in performing activities of daily living that has been present for a minimum period of 6 months.† Depending on the underlying pathology and clinical manifestation of the disease, dementia can be further grouped into Alzheimer’s disease (AD), Lewy body dementia (DBL), frontotemporal dementia (FTD),‡ and vascular dementia. Alzheimer’s disease accounts for 60% to 80% of dementia cases, and the incidence increases with the age of the patients, especially older than 65 years. It is the sixth leading cause of mortality in the United States and causes a considerable financial burden to the country. In 2012, the estimated overall care expenses accounted to US $200 billion.†

The diagnosis of AD was mainly based on the clinical manifestation of symptoms. However, advances in genetics, the development of biomarkers of neurodegeneration, and neuroimaging has lead to the incorporation of these modalities in the diagnosis of AD. It is believed that the pathophysiology of AD starts years ahead of the manifestation of clinical signs, and this discovery mandates the use of methods to detect AD earlier than conventional diagnostic tools.‡ Most available treatment modalities aim at slowing the progression of disease and controlling symptoms, and this further stresses the importance of early diagnosis of the disease.† F-FDG-PET/CT has been not only a valuable tool in tumor imaging‡–11 but also a very promising neuroimaging tool in the diagnosis of AD because it reflects resting state cerebral metabolic rates of glucose, which is an indicator of neuronal activity, and several studies have shown that cerebral metabolic alterations precede the clinical manifestation of AD symptoms. The distinct pattern of cerebral glucose metabolism also help in differentiating AD from other causes of dementia.‡ Recently, PET imaging tracers, which correlate β-amyloid deposition in the brain, have been approved by the Food and Drug Administration (FDA). The amyloid deposition in the brain can be detected years before the onset of clinical symptoms. The PET imaging tracers help in differentiating dementia syndromes, which do not have overlap of the underlying pathological process.‡ The purpose of this article was to review the role of brain PET imaging in the diagnosis of AD.

CLINICAL FEATURES, DIAGNOSIS, AND TREATMENT OF AD

The clinical manifestation of AD demonstrates a spectrum of presentations, depending on its severity. The National Institute of Neurological and Communicative Disorders and Stroke and Alzheimer’s Disease and Related Disorders Association have worked together to establish criteria to assist the clinical diagnosis of AD. The first manifestation of AD symptoms, also referred to as the preclinical phase of AD, is termed mild cognitive impairment (MCI). This entity is characterized by lower performance in 1 or more cognitive domains for the age and educational background of the patient in the absence of dementia and preservation of independence in functional abilities.‡ The National Institute of Neurological and Communicative Disorders and Stroke—Alzheimer’s Disease and Related Disorders Association classifies AD broadly as “probable AD dementia” or “possible AD.” Probable dementia is diagnosed when the patient, in addition to the criteria for all-cause dementia, has insidious onset of symptoms, clear history of cognitive worsening by report or observation, and the cognitive deficits present with either amnestic or nonamnestic presentations, in the absence of evidence of all other causes of dementia. Possible dementia of AD is considered in patients who present with an atypical course of cognitive impairment or have a mixed presentation, such as concomitant cerebrovascular disease, features of DLB or evidence of another neurological disease or nonneurological medical comorbidity, or medication use that could affect cognition.

Available treatment options for AD aim at slowing the progression of the disease and controlling symptoms. Drugs under development are aimed at targeting the pathological processes leading to AD. The approved groups of drugs for the treatment of AD include the acetylcholinesterase inhibitors, which aim at increasing the levels of acetylcholine at the sites of neurotransmission and memantine, a noncompetitive N-methyl-aspartate receptor antagonist. The common adverse effects of acetylcholinesterase inhibitors

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Dr Subramaniam was a speaker for Eli Lilly in 2013 in brain amyloid PET and received speaker fees. The remaining authors and staff in a position to control the content of this CME activity and their spouses/life partners (if any) have disclosed that they have no other financial relationships with, or financial interests in, any commercial organizations pertaining to this educational activity. Lippincott CME Institute has identified and resolved all conflicts of interest concerning this educational activity.

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ISSN: 0363-9762/14/3910-e413

Clinical Nuclear Medicine • Volume 39, Number 10, October 2014 www.nuclearmed.com | e413

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are nausea, vomiting, diarrhea, sleep disturbances, muscle cramps, weakness, bradycardia, and urinary incontinence. The adverse effects encountered with memantine include dizziness, confusion, headache, and incontinence and is used with caution in patients with renal failure or epilepsy.7

PATHOPHYSIOLOGY OF AD

Alzheimer’s disease is characterized by the accumulation of the β-amyloid peptide (Aβ) within the brain along with neurofibrillary tangles of hyperphosphorylated tau protein.15 Amyloid deposition has been described to be caused by the deposition of Aβ, a cleavage product of the amyloid precursor protein that originates from degenerating mitochondria in dystrophic neurons, and the deposition causes further disruption of axons and further deposition of amyloid.16,17 Neurofibrillary tangles caused by hyperphosphorylation of tau protein have also been identified as an important pathophysiological step in the development of AD. Studies have shown significant correlation between the concentration of phosphorylated tau protein and neurofibrillary tangles in patients with AD.16 Based on these pathophysiological processes, the measurement of certain biomarkers such as β-amyloid peptide (Aβ42), total tau, and phosphorylated tau, reflect the pathological features of AD.19 Studies have shown that these markers may be used to monitor patients with AD overtime and can also serve as a surrogate marker for treatment efficacy.20 Another biomarker of importance is the apolipoprotein E (ApoE). It has been implicated that ApoE isoforms influence clearance and deposition of Aβ. The ApoE is coded by the APOE gene, which has 3 major alleles; ε2, ε3, and ε4. Apolipoprotein E-ε4 (ApoE4) has been recognized as the strongest risk factor for sporadic AD.21

FDG BRAIN PET IMAGING

Normal Cerebral Glucose Metabolism With Aging

Cerebral glucose metabolism patterns are similar among individuals who are age matched. The mean cerebral glucose metabolism has been found to gradually decrease with age. A study by Kuhl et al22 evaluating the effects of aging on the cerebral glucose metabolism of 40 normal subjects showed that the average cerebral glucose metabolic rate at age 78 years was 26% less than at age 18 years. Within the brain, the anatomical regions that show the greatest decrease in FDG uptake with aging are the bilateral superior medial frontal, motor, anterior, and middle cingulate and bilateral parietal cortices. The superior temporal pole was found to be particularly affected. The regions that show the least changes in glucose metabolism with aging are the bilateral medial temporal lobes, putamen, pallidum, lateral thalamic nuclei, right posterior cingulate cortex, precuneus, and both sides of the occipitotemporal cortex.23

Value of FDG-PET in the Evaluation of AD

FDG-PET has been proven to be a promising modality for detecting functional brain changes in AD, identifying changes in early AD, and helping to differentiate AD from other causes of dementia. Many studies have been published evaluating the value of FDG-PET in AD for the last 3 decades. A meta-analysis including 27 studies evaluating FDG-PET in the diagnosis of AD resulted in a pooled sensitivity (SN) of 91% (95% confidence interval [CI], 86%-94%) and specificity (SP) of 86% (95% CI, 79%-91%). The analysis included 119 studies evaluating the role of different modalities in the diagnosis of AD. The results from the meta-analysis showed that FDG-PET has superior diagnostic accuracy in comparison with the other available diagnostic methods such as clinical guidelines, MRI, CT, SPECT, and biomarkers.24 Studies have also shown that FDG-PET has the potential to differentiate patients with AD from normal subjects with an SN and SP of 99% and 98%, respectively, from patients with DLB with an SN and SP of 99% and 71%, respectively, and from patients with FTD with an SN and SP of 99% and 65%, respectively.

FDG-PET in MCI

In patients with MCI, FDG-PET has been found to be better than other imaging modalities in diagnosing and predicting conversion of MCI to dementia.26 This has a significant impact on the management strategy and quality of life of affected patients because all available treatment options aim at slowing the progression of the disease and also help patients plan ahead and make important life decisions.

Studies have been published evaluating the pattern of glucose metabolism in patients with AD and MCI. Del Sole et al25 evaluated FDG-PET scans performed in 16 patients with MCI and in 14 patients with AD showing that the areas of decreased glucose metabolism involved the posterior cingulate cortex (PCC), precuneus, inferior parietal lobule, and middle temporal gyrus in patients with AD. In patients with MCI, the authors found decreased glucose metabolism only in the PCC. The areas of decreased glucose metabolism in the PCC was found to be wider in patients with AD, compared with those with MCI and extended to the precuneus. Decreased glucose metabolism in the lateral parietal cortex was found only in AD. The study has also shown that 86%, 71%, 64%, and 35% of patients with AD demonstrated decrease in the cerebral glucose metabolic rate in the PCC, temporal cortex, parietal cortex, and frontal cortex, respectively. The corresponding percentages for patients with MCI were 56%, 44%, 18%, and 0%, respectively.25 Other studies have shown changes in glucose metabolism in similar regions of the brain27,28 (Fig. 1). Another important application of FDG-PET in the evaluation of patients with MCI is its ability to predict progression to dementia. In a meta-analysis by Yuan et al29 of 24 studies involving 1112 patients, the pooled SN, SP, positive likelihood ratio, negative likelihood ratio, and odds ratio of FDG-PET to predict conversion of MCI to AD were 88.8%, 84.9%, 4.61, 0.15, and 40.15, respectively. Regions of decreased cerebral glucose metabolic rate have been observed in the temporoparietal, inferior parietal, medial temporal, and posterior cingulate cortices in patients with MCI who developed AD within a year, in comparison with normal subjects and patients with stable MCI. The progression of the disease is accompanied by a continued decrease in the glucose uptake in the regions and emergence of a new affected region involving the lateral prefrontal cortex.30–32 Mosconi et al31 evaluated 37 patients with MCI, followed them up for a year, and found that all patients who progressed to dementia showed reduced cerebral glucose metabolic rates in the inferior parietal cortex as compared with those patients who did not. Anchisi et al32 evaluated 67 patients with MCI and found that patients who converted to AD showed bilateral hypometabolism in the inferior parietal, posterior cingulate and medial temporal cortices, whereas those with stable MCI had hypometabolism in the dorsolateral frontal cortex.

FDG-PET in AD

As mentioned earlier, in patients with early AD, the areas of glucose hypometabolism have been commonly observed in the parietotemporal association cortices, posterior cingulate cortex, and the precuneus (Figs. 2 and 3). As the disease progresses, the affected regions spread to involve the frontal cortices, whereas the metabolism in the striatum, thalamus, primary sensorimotor cortices, visual cortices, and cerebellum are relatively preserved.33,34 Silverman et al35 evaluated 146 patients undergoing evaluation for dementia, of whom 97 patients were histopathologically confirmed to have AD and observed focal hypometabolism in the parietal, temporal, and/or frontal cortices or global hypometabolism. Hoffman et al36 evaluated 22 patients with
dementia, of whom 16 were found to have a pathological diagnosis of AD, and observed that bilateral temporoparietal hypometabolism had relatively high SN (93%), positive predictive value (81%), and negative predictive value (83%) as well as lower SP (63%) for the diagnosis of AD. This study proves that the finding of bilateral temporoparietal hypometabolism highly correlates with the pathological diagnosis of AD. Minoshima et al evaluated the glucose metabolism patterns in patients with autopsy-confirmed AD and found that the patients had significant metabolic reductions in the lateral parietal, temporal, frontal, and posterior cingulate cortices. They also observed either mild or insignificant reduction in the glucose metabolism in the primary visual, sensorimotor cortices and subcortical structures. Ossenkoppele et al compared the SUV ratio (SUVr) obtained with the cerebellar grey matter as the reference tissue and observed that in patients with AD at baseline, there was a decrease in SUVr in the parietal, posterior cingulate and temporal cortices. At a mean interval follow-up of 2.5 years, the authors observed that there was a decrease in SUVr from baseline in the frontal, parietal, and lateral temporal lobes over time in patients with AD.

FDG-PET in Differentiating AD From Other Types of Dementias

An important application of FDG-PET in clinical practice in the context of dementia is its ability to differentiate AD from other causes of dementia. Many studies have been performed to help clinicians differentiate between the types of dementias. Gilman et al evaluated 25 patients with AD, 20 with DLB, and 19 normal control subjects and found that the FDG-PET studies showed significantly lower cerebral glucose metabolic rates in the visual cortex (Brodmann areas 17, 18, and 19) in patients with DLB compared with patients with AD. They also observed no significant difference in the glucose metabolism in the PCC, superior parietal lobe, lateral temporal lobe, and prefrontal region between the 2 groups. However, a study by Lim et al evaluating 14 patients with DLB and 10 patients with AD has shown that hypometabolism in the lateral occipital cortex had the highest SN (88%) and the relative preservation of the mid or posterior cingulate gyrus had the highest SP (100%) in differentiating DLB from AD. Minoshima et al have shown that significant cerebral glucose metabolic reduction in the occipital cortex, especially in the primary visual cortex, distinguished DLB from AD with an SN and SP of 90% and 80%, respectively (Fig. 4). Clinically, it is important to differentiate patients with DLB from patients with AD because patients with DLB can be severely dopaminergic deficient and can adversely react severely to neuroleptic treatment.

Foster et al evaluated the glucose metabolism patterns in patients with pathologically confirmed AD from those with FTD. The authors found glucose hypometabolism in the frontal, anterior cingulate and anterior temporal regions more frequently in patients with FTD, whereas patients with AD demonstrated hypometabolism in the temporoparietal and posterior cingulate regions, more commonly. The authors found a mean diagnostic accuracy of 84.8% by transaxial FDG-PET imaging and 89.2% by stereotactic surface projection of the FDG-PET images (Fig. 5). Clinically, it is important to differentiate patients with FTD from patients with AD because patients with...
FTD can have serious adverse effects if treated with anticholinesterase inhibitors.

AMYLOID BRAIN PET IMAGING

Although the etiology of AD has not been fully established, there is evidence to suggest that the amyloid-β peptide plays an important role in AD pathogenesis.\(^4\) Accumulation of Aβ fibrils in the form of amyloid plaques is a neuropathological hallmark for autopsy-based diagnosis confirmation of dementia caused by AD,\(^4\) and more importantly, Aβ deposition is thought to precede cognitive symptoms in AD and is therefore a potential preclinical marker of disease.\(^4\) There have been different approaches to noninvasively visualize amyloid deposition in human brains with amyloid PET radiotracers.

PET Amyloid Radiopharmaceuticals

Typically, amyloid imaging agents bind to insoluble fibrillar forms of Aβ 40 and Aβ 42 deposits, which are a major component of compact neuritic plaques and vascular deposits. PET amyloid-β imaging agent could facilitate the clinical evaluation of late-life cognitive impairment by providing an objective measure for AD pathology. PET imaging probes of Aβ plaques have been extensively developed during the last decade.

\(^{11}\)C-Pittsburgh Compound B (PiB) was the first amyloid imaging PET agent used in humans subjects in 2002.\(^4\) In April 2012, the US FDA approved the first Aβ imaging PET probe, \(^{18}\)F-florbetapir\(^6\) (Amyvid, Eli Lilly and Company), to identify Aβ plaque accumulation in patients with suspected AD. Subsequently, another amyloid PET radiopharmaceutical derived \(^{11}\)C-PiB, flutemetamol (Vizamyl, GE Healthcare)\(^7\) and more recently a third amyloid imaging probe, \(^{18}\)F-florbetaben injection\(^8\) (Neuraceq, Piramal Imaging) had been approved by the FDA. Many other amyloid ligands are available for research use, such as the benzofuranes \(^{11}\)C-AZD-2184 and \(^{18}\)F-AZD-4694,\(^9\) benzoxazole \(^{11}\)C-BF-227,\(^10\) as well as stilbene compounds \(^{11}\)C-SB,\(^11\) and naphthol \(^{18}\)F-FDDNP.\(^12\) The following discussion will focus on FDA-approved amyloid PET radiotracers.

\(^{11}\)C-PiB \([\text{N-methyl-}^{11}\text{C}]2-(4-methylaminophenyl)-6-hydroxybenzothiazole]\)

\(^{11}\)C-PiB is a derivative of a fluorescent amyloid dye, thioflavin T, and has been found to possess high affinity and high SP for fibrillar Aβ.\(^4,5\)\(^3\) It was developed by Chet Mathis and William Klunk at the University of Pittsburgh and was first used in human research studies in 2002 in collaboration with Uppsala University, Sweden. The compound was named Pittsburgh Compound B (\(^{11}\)C-PiB). The initial human study of \(^{11}\)C-PiB was expanded to include 16 subjects with AD and 9 cognitively healthy controls, which was published in 2004.\(^5\) Patients with AD were found to show significantly higher \(^{11}\)C-PiB retention in the frontal cortex (1.94-fold), parietal cortex (1.71-fold), corpus striatum (1.76-fold), temporal cortex (1.52-fold), and occipital cortex (1.52-fold). Since that study, \(^{11}\)C-PiB imaging has been rapidly spread to academic centers worldwide and widely used in many research studies. PiB compound is labeled with \(^{11}\)C, with short...
half-life of only 20 minutes, limiting its use to PET centers equipped with an on-site cyclotron and \(^{11}\)C radiochemistry expertise. To overcome these limitations, \(^{18}\)F-labeled A\(_{\text{A}}\) tracers, with longer half-life of 110 minutes, have been developed and found to successfully correlate with \(^{11}\)C-PiB results, providing a reliable assessment of brain amyloid with a single scan of 15- to 20-minute duration.

**\(^{18}\)F-Florbetapir (\(^{18}\)F-AV-45; Amyvid)**

\(^{18}\)F-Florbetapir was the first amyloid imaging agent approved by the FDA in April 2012.\(^{54}\) It is a stilbene derivative, first synthesized by Kung et al at the University of Pennsylvania. It has rapid reversible binding characteristics, allowing scanning to commence 45 to 50 minutes after injection, similar to \(^{11}\)C-PiB. There are a number of studies evaluating the efficacy of florbetapir in detecting A\(_{\text{A}}\) pathology. In a multicenter study of 59 patients with different levels of cognitive function, florbetapir showed an SN and SP of 92% and 100%, respectively, in detecting amyloid plaques.\(^{44,46,54-56}\) The accumulation of amyloid detected by florbetapir PET imaging significantly correlated with postmortem amyloid assessment.\(^{56}\) Rowe et al\(^{48}\) also reported florbetapir binding in areas matching the postmortem A\(_{\text{A}}\) plaques distribution. Brain retention of \(^{18}\)F-florbetapir is also highly correlated with \(^{11}\)C-PiB (\(r = 0.97\)).

**\(^{18}\)F-flutemetamol (also known as GE067) (Vizamyl, GE Healthcare)**

Flutemetamol is another \(^{18}\)F-labeled amyloid imaging approved by the FDA in 2013. \(^{18}\)F-flutemetamol has demonstrated dosimetry comparable with other \(^{18}\)F-labeled radiopharmaceuticals and performed similarly to \(^{11}\)C-PiB with high correlation (\(r = 0.91\)).\(^{46,54,56-58}\) Studies have provided additional data supporting the concordance between in vivo \(^{18}\)F-flutemetamol imaging and histopathology.\(^{59}\)

**AMYLOID PET IMAGING IN AD**

Studies suggest that the visual inspection of PET imaging amyloid scan shows a typical regional brain distribution in patients with AD, which seems to replicate the sequence of A\(_{\text{A}}\) deposition found at autopsy,\(^{46,54,56,59,60}\) with initial deposition in the precuneus, orbitofrontal cortex, and inferior temporal, posterior cingulate gyrus, followed in time by the remaining prefrontal cortex as well as lateral temporal and parietal cortices. There is relative sparing of the sensorimotor, occipital, and medial temporal regions.

A negative amyloid PET scan result (Fig. 6) indicates a reduced likelihood that cognitive impairment is caused by AD, but a positive scan result (Fig. 7) does not establish a diagnosis of AD. However, amyloid PET imaging must be performed in a setting of a clinically suspected AD because there is a relative low SP. Klunk et al conducted a review of 15 studies with a total of 341 patients with

\[\text{FIGURE 3. Alzheimer’s disease:} \quad \text{\(^{18}\)F-FDG brain PET/CT study of a 81-year-old man with cognitive decline during the past few months. The left and right lateral (A) and left and right medial (B) 3-dimensional stereotactic surface projection images of FDG hypometabolism demonstrate glucose hypometabolism involving the bilateral frontal, temporal, and parietal cortices (compared with a normal brain database, Mm software 6.2. Light blue, \(-1\) SD; dark blue, \(-2\) SDs; purple, \(-3\)SDs), with sparing of the sensorimotor and occipital cortices, features consistent with AD.} \]
AD and 651 subjects with normal cognitive performance, concluding that 96% of the patients with clinically suspected AD had amyloid imaging positive result, with an SP of 76%. The degree of cortical binding of amyloid agent in patients with AD is highly variable and does not correlate with clinical measures of cognitive impairment severity. This has been shown in longitudinal studies assessing for disease progression demonstrating stable PiB retention after 2 years of follow-up in a whole spectrum of cognitive stages, from cognitively unimpaired individuals to patients with AD dementia. A accumulation is a slow process, and the evidence suggests that it remains constant in the preclinical and prodromal stages of the disease.

There have been efforts to evaluate the impact of amyloid PET in clinical management of patients with AD. Grundman et al assessed the impact of florbetapir imaging in 229 patients, reporting change in the diagnosis 54.6% of patients (CI, 48.1%–60.9%) after amyloid imaging scan with 18F-florbetapir, and the diagnostic confidence was increased by an average of 21.6% (CI, 18.3%–24.8%). Thus, the incorporation of amyloid PET into the management of patients with AD may be valuable, offering the possibility of monitoring and assessing the effectiveness of pharmaceutical therapeutic agents directed against Aβ levels. Rinne et al conducted a phase II study (n = 19 patients), where PiB PET imaging was used to investigate whether bapineuzumab, a monoclonal antiamyloid antibody, would reduce cortical fibrillar Aβ load in patients with AD. The difference in mean PiB retention ratio changes between the bapineuzumab and the placebo group was −0.24 (P < 0.003). The differences in the individual regions between the 2 groups were similar. Ostrowitzki et al studied the effectiveness of gantenerumab treatment using PiB PET imaging (n = 16 patients) and found that using 60 mg of gantenerumab decreased the Aβ deposition by 15.6% (95% CI, −24.0 to 11.6%), whereas using 200 mg of gantenerumab decreased the deposition by 35.7% (95% CI, −63.5% to −7.9%). These studies illustrate the potential value of amyloid imaging in assessing the effectiveness of new treatment options against Aβ in clinical trials as well as monitoring the effect of treatment in clinical practice.

Amyloid PET Imaging in Mild Cognitive Disorder

Approximately 25% to 35% of elderly subjects with adequate performance in cognitive tests demonstrate high cortical radiotracer amyloid imaging retention in the prefrontal, posterior cingulate and precuneus regions. These findings are concordant with postmortem reports, which show that approximately 25% of non-demented subjects older than 75 years have Aβ plaques. In nondemented subjects, the presence of Aβ deposition might reflect a preclinical stage of AD.

Unlike AD, there is correlation between amyloid imaging binding, using 11C-PiB, and the degree of memory impairment in nondemented subjects, and approximately 50% to 60% of individuals with MCI will progress to AD during 3 to 5 years of follow-up. There is evidence that a positive amyloid PET scan result in patients with MCI will help in predicting conversion to AD and, thus, can potentially help to identify patients who will benefit from specific therapies. This was assessed by Jack et al in 218 patients who concluded that amyloid positive scan results were associated with a significantly higher chance to progress to AD, with a hazard ratio of 3.2 (P = 0.004).

FIGURE 4. Lewy body dementia: 18F-FDG brain PET/CT study of a 43-year-old woman with memory loss and visuospatial dysfunction. The left and right lateral (A) and left and right medial (B) 3-dimensional stereotactic surface projection images of FDG hypometabolism demonstrate glucose hypometabolism involving the bilateral occipital and parietal cortices, extending to the temporal cortex (compared with a normal brain database, Mim software 6.2. Blue, −2 SDs; purple, −3 SDs), features suggestive of Lewy body dementia.
20% of subjects with of $^{11}$C-PiB–negative MCI progress to other types of dementia, such as Lewy bodies dementia or frontotemporal dementia.

**Appropriate Use Criteria for Brain Amyloid Imaging with PET in AD**

Criteria for appropriate clinical use are needed to avoid potential patient harm if the amyloid imaging scans are performed in inappropriate circumstances or are poorly read or if the significance of the results is not correctly applied to the clinical context.

The Society of Nuclear Medicine and Molecular Imaging and the Alzheimer’s Association Amyloid Imaging Taskforce have jointly developed consensus recommendations for the appropriate use of brain amyloid imaging to aid in the diagnosis of people with suspected AD. The Amyloid Imaging Taskforce concluded that amyloid imaging could potentially be helpful in the diagnosis of...
A or cerebral amyloidosis. Therefore, the potential clinical use of amyloid PET requires careful consideration in the proper clinical setting.

Another limitation is that a positive amyloid scan result can also be seen not only in AD but also in other medical conditions, such as dementia with Lewy bodies,\(^\text{46,54,56,70,71}\) or cerebral amyloid angiopathy.\(^\text{46,54,55,71,72}\) Thus, it is important to emphasize that amyloid positivity does not establish differential diagnosis between AD and A\(\beta\) disorders. Furthermore, amyloid PET would not add any useful information in differentiating disorders that are not associated with A\(\beta\), such as frontotemporal dementia syndromes, or in detecting rare forms of AD in which ligand binding is greatly reduced because of unusual forms of A\(\beta\).\(^\text{46,54,56,72,73}\)

**CONCLUSIONS**

FDG-PET and amyloid PET imaging are valuable in the assessment of patients with dementia. The characteristic spatial distribution of glucose metabolism on brain FDG-PET can help in differentiating AD from other causes of dementia such as frontotemporal dementia and dementia of Lewy body. A negative amyloid brain PET result is useful in excluding significant amyloid deposition and thus AD in appropriate clinical setting.

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**CME EXAMINATION**

Please mark your answers on the ANSWER SHEET.

**CME Learning Objectives:**

After completing this journal-based CME activity, participants will be able to:

- Identify patterns of FDG spatial distribution in brain PET of patients with MCI, AD, frontotemporal dementia, and dementia of Lewy Body.
- Identify the value and learn the appropriateness criteria for amyloid brain PET.

1. In Figure 1A and B, you are shown 3-dimensional stereotactic surface projection medial and lateral images of FDG metabolism (1A) and FDG hypometabolism (1B, z score, compared with a normal brain database, GE advantage workstation software 4.2, Milwaukee, WI. Yellow, −2 SDs; red, −3 SDs), respectively, of a 51-year-old woman who presented with a history of progressive nonfluent aphasia during the past 9 months. What condition does the glucose metabolism pattern most likely demonstrate?

A. MCI
B. Alzheimer dementia
C. Lewy body dementia
D. Frontotemporal dementia

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2. In Figure 2A and B, you are shown axial FDG-PET and fused FDG-PET/CT images of a 87-year-old woman who presented with a recent history of cognitive decline. What type of dementia does the glucose metabolism pattern most likely demonstrate?

A. MCI  
B. Alzheimer dementia  
C. Lewy body dementia  
D. Frontotemporal dementia

Please see the following references for further study:

3. Which of the following diagnostic modalities has the highest SN and SP to diagnose AD?

A. Clinical guidelines  
B. SPECT  
C. FDG-PET  
D. Amyloid β1-42 in combination with tau

Please see the following references for further study:

4. Which of the following ApoE isoforms has been recognized as the strongest risk factor in sporadic AD?

A. ApoE-ε4  
B. ApoE-ε2  
C. ApoE-ε3  
D. ApoE-ε1

Please see the following references for further study:

5. In differentiating MCI from AD using FDG-PET imaging, which of the following structures is usually preserved in MCI in comparison with AD?

A. Posterior cingulate cortex  
B. Temporal cortex  
C. Parietal cortex  
D. Frontal cortex
Please see the following references for further study:


*6. With progression of disease, which of the following structures exhibit relatively well-preserved glucose metabolism in patients with AD?*

A. Parietotemporal cortices  
B. Precuneus  
C. Visual cortex  
D. Posterior cingulate cortex

Please see the following references for further study:


*7. In differentiating dementia with Lewy bodies from Alzheimer dementia using FDG-PET, which of the following structures shows significant difference in the glucose metabolism pattern between the 2 disease groups?*

A. Posterior cingulate cortex  
B. Superior parietal lobe  
C. Lateral temporal lobe  
D. Visual cortex

Please see the following references for further study:


*8. Which was the first amyloid PET imaging aging used in humans?*

A. 18F-AZD-4694  
B. 11C-PiB  
C. 18F-florbetapir  
D. 18F-florbetaben

Please see the following references for further study:


*9. Which of the following patients will not be an appropriate candidate for amyloid PET imaging evaluation?*

A. A 77-year-old man with persistent unexplained MCI  
B. A 70-year-old man, asymptomatic, with family history of AD.  
C. A 68-year-old woman with unclear clinical presentation for AD, with a positive test result for possible AD.  
D. A 55-year-old man patient with progressive cognitive impairment.

Please see the following references for further study:


*10. A positive amyloid PET imaging can be seen in the following cases, except*

A. Dementia with Lewy bodies  
B. AD  
C. Frontotemporal dementia  
D. Cerebral amyloid angiopathy
Please see the following references for further study:

*11. Which of the following PET radiotracer is not$^{18}$F labeled?*

A. PiB  
B. Flumetamol  
C. Florbetapir  
D. Florbetaben

Please see the following references for further study:

Answer sheet can be found online in the October 2014 issue of *Clinical Nuclear Medicine* located at www.nuclearmed.com. All recipients of print subscription already have access to the online version of the journal. Please contact customerservice@lww.com, or call customer service at 800-638-3030 or 301-223-2300 for help with registering and logging in to the website. Or you can visit [http://CME.LWW.com](http://CME.LWW.com) for more information about this educational offering and to complete the CME activity online.