The degree of intactness of human cognitive functioning for a given individual spans a wide spectrum, ranging from normal to severely demented. The differential diagnosis for the causes of impairment along that spectrum is also wide, and often difficult to distinguish clinically, which has led to an increasing role for neuroimaging tools in that evaluation. The most frequent causes of dementia are neurodegenerative disorders, Alzheimer’s disease being the most prevalent among them, and they produce significant alterations in brain metabolism, with devastating neuropathologic, clinical, social, and economic consequences. These alterations are detectable through positron emission tomography (PET), even in their earliest stages. The most commonly performed PET studies of the brain are performed with $^{18}$F-fluorodeoxyglucose as the imaged radiopharmaceutical. Such scans have demonstrated diagnostic and prognostic utility for clinicians evaluating patients with cognitive impairment and in distinguishing among primary neurodegenerative disorders and other etiologies contributing to cognitive decline. In addition to focusing on the effects on cerebral metabolism examined with $^{18}$F-fluorodeoxyglucose PET, some other changes occurring in the brains of cognitively impaired patients assessable with other radiotracers will be considered. As preventive and disease-modifying treatments are developed, early detection of accurately diagnosed disease processes facilitated by the use of PET has the potential to substantially impact on the enormous human toll exacted by these diseases.

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Among all applications of positron emission tomography (PET) of the brain in current clinical practice, evaluating changes in cognitive abilities is the one for which the greatest demand currently exists. The degree of intactness of human cognitive functioning spans a wide spectrum, including (1) normal for a given individual; (2) subjective perception of impairment that is not objectively documented by neuropsychologic tests but which can represent the earliest stages of decline; (3) minimally impaired states with neuropsychologic functioning diminished slightly below the mean level of normal young performance (sometimes categorized as “age-associated” or “age-consistent”); (4) an intermediate state often referred to as “mild cognitive impairment” (MCI); and (5) states of dementia of mild, moderate, or severe degree. Although some people experiencing milder forms of impairment are on a path of progressive decline, others will have stable or reversible forms of impairment, and the differential diagnosis for the associated causes is wide at every level, with the actual cause often difficult to clinically discern.

Dementia affects approximately 24 million people worldwide, with about 5 million new cases occurring annually. The most common cause of decline that eventually leads to dementia, and the best-studied, is Alzheimer’s disease (AD). This condition occurs in 10% to 15% of people older than 65 years of age, and 40% to 50% of all those older than 85 years of age. The disease is present in approximately two-thirds of cases of dementia. It is estimated to currently afflict 5.2 million people at a cost to society of $148 billion in the United States alone, a number projected to increase to 11 to 16 million people by 2050, as the aging of the baby boomers expands our geriatric population. Other causes of cognitive decline include dementia with Lewy bodies, cerebrovascular disease, frontotemporal dementia, Creutzfeld–Jakob disease, HIV-associated dementia, neurosyphilis, Parkinson’s dementia, normal pressure hydrocephalus, and dementias caused by exposure to toxic substances (heavy metals, alco-
hol, other drugs), metabolic abnormalities, or psychiatric disorders.

Limitations of Conventional Clinical Evaluation

A definite diagnosis of AD can be made only by histopathologic examination of brain tissue. The identification and differential diagnosis of AD is especially challenging in its early stages, partly because of the difficulty in distinguishing it from the mild decline in memory that can occur with normal aging and from mild cognitive manifestations of other neuropsychiatric conditions, such as depression, as well as other causes of dementia. False-positive and false-negative diagnoses with respect to presence of AD are not uncommonly made, even by physicians experienced with the evaluation of dementia. Moreover, clinical assessment frequently involves multiple examinations and laboratory tests over the course of months or years.

There is greater need for early accurate diagnosis, now that several medications for the treatment of mild-to-moderate AD are available. Most are cholinesterase inhibitors, used to delay cognitive decline. More recently, memantine (Namenda; Forest Laboratories, Inc, New York, NY) was approved by the United States Food and Drug Administration for treating patients with moderately advanced disease. Patients with neurodegenerative disease have the most to gain from therapy that intervenes as early as possible in the course of inexorably progressive irreversible damage to brain tissue, and controlled clinical trials have demonstrated that cholinesterase inhibitors can improve, or delay the decrease in, memory and other cognitive functions in mild or moderately affected AD patients. Treatments can reduce by more than half the proportion of patients requiring nursing home placement over a given period of time. Cholinergic and other agents also have beneficial effects with respect to reducing behavioral problems, improving patients’ functional abilities, and decreasing caregiver burden.

Studies examining the long-term effects of cholinesterase inhibitors indicate that drug treatment produces an average delay in cognitive decline in AD patients of 9-12 months and a delay in the need for institutionalization of 18 months on average. Moreover, delaying the implementation of therapy by as little as 6 months, in addition to carrying the inherent adverse consequence of depriving the patient of the short-term advantages of potentially enhanced cognitive and functional abilities during that time, may also have long-term consequences. Because early treatment interventions are able to keep patients at greater levels of functioning longer, and future innovative therapies may be able to further delay the onset of dementia and slow its progression, or potentially reverse it when intercepted at a sufficiently early stage, it is becoming increasingly important to accurately diagnose dementia as early as possible.

Few studies have specifically addressed the question of clinical detection of very mild disease, however, particularly in comparison with the criterion standard of histopathologic diagnosis. In the report of the Quality Standards Subcommittee of the American Academy of Neurology, 3 “Class I” studies were identified in which the diagnostic value of clinical assessment could be meaningfully measured (Class I indicates “a well designed prospective study in a broad spectrum of persons with the suspected condition, using a ‘gold standard’ for case definition, and enabling the assessment of appropriate tests of diagnostic accuracy”), and only one focused on evaluating dementia at a relatively early stage. To be included in that investigation, patients were required to have had onset of dementia symptoms within 1 year of entry. All of the 134 patients evaluated underwent a complete standardized diagnostic workup composed of a comprehensive medical history and physical, neurological examination, neuropsychologic testing, laboratory tests, and structural neuroimaging, in addition to an average of 3 additional years of clinical follow-up with repeated testing. Sensitivity of this assessment for AD was 83-85%, whereas specificity was 50-55%, yielding an overall accuracy of 69%.

An impediment to accurately identifying presence of disease is reliance on standardized cut-off scores on mental status or cognitive screening tests. Highly educated individuals who have suffered cognitive decline may show normal performance on cognitive examinations, whereas people with less education may appear to have cognitive impairment or dementia when they actually have not declined. Test scores are most useful as a quantitative baseline against which to compare future assessments, rather than as diagnostic markers. A particular problem in the early stages of decline is that clinicians often have difficulty distinguishing complaints of the “worried well” (ie, normal aging) from those of patients who have an underlying brain disorder that will result in progressive cognitive decline. Subjective complaints can indicate the presence of mood disorders or early dementia, and in any event, should be taken seriously.

FDG-PET Studies of Normal Adult Brain and Healthy Aging

Clinical PET studies of the brain are performed most commonly with the use of fluorodeoxyglucose (FDG). The resulting scans are typically interpreted qualitatively, by visual analysis, although quantitative software tools are becoming increasingly used as adjuncts to interpretation. In either case, the relative distribution of FDG throughout the patient’s brain is examined, and compared with the distribution expected for a normal subject.

When quantification is performed, it may be expressed as regional concentrations of measured radioactivity normalized to some internal reference standard, for example, a reference region of the brain, the whole brain activity, or the average whole-body concentration before excretion and decay-corrected to the actual time of imaging (ie, standardized uptake value, or SUV). Those results, often termed “semi-quantitative,” turn out to be adequate for most clinical applications, as well as for many research applications. In contrast, “absolute quantitative” values are derived from biologically
based mathematical models that reflect the partitioning of radioactivity into compartments that can reflect both physiological boundaries (eg, the vascular space, the blood–brain barrier, the plasma membrane of neurons) and biochemical processes (enzymatic anabolism and degradation, transport molecules). These models necessarily represent substantial simplifications of the actual biological environment but nevertheless have proven capable of yielding quantitative estimates in good agreement with similar measures obtained by more invasive methods.

In the case of FDG studies, the biological parameter that is being estimated is the rate of regional glucose utilization, based on a method described by Sokoloff and his colleagues, originally developed with 14C-labeled 2-deoxyglucose. Early measures of regional glucose utilization rates in the human brain yielded estimates of global cerebral metabolism of approximately 5.5 mg glucose/min/100 g, which ranged from 3.6 to 5.2 mg glucose/min/100 g in white matter to 5.8–10.3 mg glucose/min/100 g in gray matter tissues. Regional values that have been more recently published reflecting measurements using instruments and techniques with improved imaging capabilities, are in substantial agreement with these initially reported values. A recent study by Yamaji and coworkers is notable for having obtained regional SUV measurements of cortex in the same group of subjects in whom absolute quantitative values were obtained, with remarkably close correspondence of the 2 types of measures observed in healthy brain. Frontal cortex was calculated to use 7.9 mg of glucose per minute per 100 g of brain tissue and had an SUV of 7.7; sensorimotor cortex demonstrated values of 8.1 and 7.8 for corresponding absolute and SUV determinations, parietal cortex 7.8 and 7.7, temporal cortex 7.1 and 7.0, and occipital cortex 7.8 and 7.7, respectively.

Cerebral blood flow can be measured with absolute quantification or semiquantitatively using the tracer H215O with PET. The rate of blood flow is normally tightly coupled with local metabolic needs of healthy brain tissue, through vasoconstrictive–vasodilatory autoregulation of blood supply. Thus, within a vascular territory, measures of cerebral blood flow and glucose metabolic rate co-vary nearly linearly. Between different vascular territories, however, different constants of proportionality can pertain. For example, because most of the lateral neocortex is supplied by the middle cerebral artery branch of the carotid circulation, the pattern of distribution of H215O closely parallels that of the metabolic tracer FDG throughout most of the cortical surface. However, dissociations can occur. The cerebellum, despite its lower metabolic activity relative to neocortex, is more readily perfused, being supplied by arterial branches of the vertebrobasilar circulation. Also, in certain pathologic circumstances (eg, cerebrovascular disease), the normal coupling between metabolism and perfusion can be disturbed, such that a consistent relationship may not exist even within a vascular territory.

With either tracer, an important issue is determining how PET-based measurements of normal brain function can be expected to change during the course of healthy aging. Effects of normal aging on adult brain function have been examined with PET. In a study of 57 healthy adults ranging in age from 19 to 50 years, the most significant age-related decline in cerebral blood flow was found in the mesial frontal cortex, encompassing the anterior cingulate cortex, and extending rostrally into the supplementary motor area. In an independent study of 27 healthy adults ranging from 19 to 76 years of age, the most significant age-related decline was found in the medial orbitofrontal cortex, and this was the only regional effect to remain significant after correction for partial volume effects of cerebral atrophy. Likewise, measures of metabolism using FDG have also identified an age-related decline in healthy adults, most consistently in frontal cortex. Nevertheless, as previously reviewed, studies of carefully selected subjects find declines to be minimal in glucose metabolism throughout most of the brain in normal aging.

**From Normal to Mild Decline in Cognition to Dementia**

As minimal gradual changes are expected with healthy aging, with respect to both cognitive abilities and cerebral metabolism, evaluating patients in the earliest stages of potentially pathologic change requires initiating that process at the time that decline first becomes evident. The decline may be noted by a physician with whom the patient has an ongoing relationship, by a family member or other close contact of the patient, or by the history provided by a patient who is deemed to be reliable. The key concept is that there has been mild decline in cognition (MDC), that is, a decline relative to the patient’s own normal level of performance, greater or faster than would be expected for normal aging.

In patients who are high-functioning from the start, MDC may occur at a much earlier point in time (often several years earlier) than the point at which the patient would meet criteria for a specific diagnostic category, such as mild cognitive impairment (MCI), as impairment is defined at least in part by how the patient performs relative to a normal group (Fig. 1). A limitation with impairment-based categories is that even when the comparative normal group is adjusted for age and educational level, there is a wide range of baseline abilities among say, 65-year-old college graduates, varying with factors such as premorbid ability, further educational achievement, differences in decades of professional, and social activity, etc.

How accurately can FDG-PET be used in the evaluation of nondemented patients who are in the earliest stages of cognitive decline? Certain patterns of regional cerebral metabolism are predictive of future cognitive decline (Fig. 2), and it is clear that metabolic changes associated with early AD, such as decreases in posterior cingulate and associative cortical activity, can be detected with PET, even before the symptomatic manifestations of the disease become evident. Overall accuracies achieved with use of FDG-PET have been nearly as high in very mildly affected patients as in demented patients. PET may be especially valuable in this clinical
setting, considering the difficulty of distinguishing these patients from those with mild memory loss attributed to normal aging.

FDG-PET may serve explicitly as a prognostic tool, to determine likelihood of deterioration of mental status in the period following the time of scanning. Relative hypometabolism of associative cortex can be accurately used to predict whether cognitive decline will occur at a rate faster than would be expected for normal aging, over the several years after a PET evaluation.42,45 Moreover, the magnitude of decline during a 2-year period, for some standardized measures of memory, correlates with the initial degree of hypometabolism of inferior parietal, superior temporal, and posterior cingulate cortical regions.46 As cognitive impairment caused by a neurodegenerative disease progresses, associated progression of regions of hypometabolism also occurs. Physicians examining brain PET scans for dementia prognosis in cognitively impaired patients with depression and thyroid disease should interpret positive scans with caution, however, because of the potentially confounding effects of those conditions on regional brain metabolism.44 For example, Bench and coworkers reported that the metabolism of certain cerebral regions are negatively correlated with mood symptoms and severity of psychomotor slowing in depression47,

Figure 1 Longitudinal visual and quantitative PET evaluations of an initially clinically “normal” subject with MDC. The first 3 columns of images represent transaxial slices through the brain, displayed from superior (parietal cortex posteriorly) to inferior (temporal cortex posteriorly) levels. Right side of image is left side of brain. PET data are displayed in an inverse linear gray scale (left color bar underneath brain images). In the fourth column, color denotes standardized volumes of interest quantifying FDG activity in parietotemporal cortex (2-dimensional color scale underneath brain images; the y-axis denotes number of standard deviations (SD), the x-axis denotes magnitude of difference from mean of asymptomatic controls.) Top row, FDG-PET scan of high-functioning 71-year-old man, who noted mild decline in his own cognition, but was considered “normal” both by general clinical evaluation and formal neuropsychologic testing. Scan reveals mild metabolic asymmetry with left temporal cortex (arrow in rightmost plane) lower than right. Left parietotemporal cortex was quantified as 3 SD and 9% below normal. Posterior cingulate cortex was also quantified as decreasing 3 SD below normal (not shown) at this point, and scan was interpreted as concerning for incipient dementia process. Middle row, Progression of posterior hypometabolism (arrows in middle and right planes). Left parietotemporal cortex was quantified as 4 SD and 12% below normal. Patient now met criteria for MCI on formal neuropsychologic testing. Bottom row, Further progression of posterior hypometabolism (arrows in left, middle, and right planes). Left parietotemporal cortex was quantified as 5 SD and 16% below normal. Patient now met criteria for borderline MCI/dementia.
these regions were found to include the inferior parietal and superior temporal cortex, areas also affected in AD. With respect to examining the added prognostic value of PET beyond that achieved with conventional clinical assessment, it has been found that among patients having clinical working diagnoses presuming progressive etiologies for their cognitive problems, those whose PET scans were nevertheless negative were 12 times more likely to remain cognitively stable than those who did not decline at the follow-up. This baseline metabolic reduction predicted decline to MCI with 83% sensitivity and 85% specificity. Although reductions in medial temporal lobe volumes do correlate with the decline from NL to MCI, and can be predictive of future dementia, this metabolic decline occurred in excess of atrophic changes. After correcting for atrophy, the decliners still showed 11% decreased entorhinal cortical metabolism as compared with the nondecliners, and the predictive accuracy was almost the same, at 80%, analogously to significant metabolic reductions which have been demonstrated to persist in MCI and AD after atrophy correction.

In addition to alterations in brain metabolism generally identifiable in MCI patients, particularly in medial temporal areas, specific regions of hypometabolism have been shown to be predictive of MCI conversion to probable Alzheimer’s disease, most consistently in parietal, temporal and posterior cingulate cortex. Predictive accuracies of these patterns have been high, typically ranging from 80% to 100%.

Genetic background concomitantly affects risk for declining from a state of normal cognitive function to developing AD, and patterns of brain glucose metabolism. In particular, the epsilon 4 allele of the apolipoprotein E gene is associated with a significantly increased risk of developing AD of senile onset; overall, elderly individuals with the 4/3 or 4/4 genotype are more than twice as likely to develop AD at any given age compared with individuals with the 3/3 genotype. FDG-PET studies have linked the epsilon 4 allele to hypometabolism in posterior cingulate, parietal, and temporal cortex, and have identified greater metabolic asymmetry in nondemented relatives of individuals with probable AD. Furthermore, significant metabolic decline in these regions has been longitudinally observed in those who have inherited the epsilon 4 allele as measured by repeating PET in the same subjects over a two-year interval, and decreased posterior...
cingulate metabolism is observable in subjects as young as in their twenties who have inherited the allele.67,68

**Diagnosis of Dementing Illnesses**

A wide variety of neurodegenerative diseases has been demonstrated to produce significant alterations in brain function detectable with PET. Distinguishing these alterations with PET has served as a subject of considerable investigation for three decades,69-72 and continues to be actively studied and extensively reviewed.44,73-75 Many thousands of patients with clinically diagnosed—and, in some cases, histopathologically confirmed—causes of dementia from many independent laboratories have been studied using PET measures of cerebral blood flow, glucose metabolism, oxygen utilization, amyloid and other microstructural imaging agents, as well as of neurotransmitter receptors, transporters, and enzyme systems. The best-studied application of this type is the use of FDG-PET to evaluate for the presence of AD. Sensitivity of FDG-PET in this context has been consistently high, indicating that by the time a patient presents with a neurodegenerative dementia, substantial alteration of cortical metabolic function has occurred (Fig. 3). The associated decreases in glucose metabolism in certain brain areas are readily detectable on FDG images, and identification of the particular brain areas of involvement can assist with the differential diagnosis of dementia.44 For example, in contrast to the relatively preserved occipital metabolism seen even in late stages of AD (Fig. 3, bottom row), occipital cortex is not preferentially spared by the dementia of Parkinson’s disease, or Dementia with Lewy Bodies (Fig. 4); similarly, subcortical structures preserved in AD are not preferentially spared in vascular dementia. Other neurodegenerative diseases with well-established characteristic patterns of hypometabolism (particularly in the less advanced stages of dementia, when assistance with accurate diagnosis is most needed) include frontotemporal dementia (prefrontal and/or anterior temporal cortex affected out of proportion to other cortical areas), and Huntington’s disease (caudate and then putamen markedly affected at a time that cortical metabolism is relatively intact).

Most studies of the diagnostic accuracy of brain imaging in dementia have been based upon comparison of PET findings with clinical assessments performed near the time of PET. The ability of that approach to determine diagnostic accuracy is limited by the fact that clinical diagnosis can be inaccurate, particularly for patients presenting in the earlier stages of disease—a time when the opportunity for effective therapy, and for meaningful planning, is greatest. Studies comparing neuropathologic examinations with imaging are thus most informative in assessing the diagnostic value of PET. In a pooled analysis10 of 3 previously published studies,77-79 histopathologically confirmed sensitivity and specificity of PET for detecting the presence of AD were 92% and 71%, respectively. In the largest single-institution series published up to that point, Hoffman and coworkers80 found sensitivity and specificity of PET for Alzheimer’s disease to occur in the range of 88% to 93% and 63% to 67%, respectively.

In a subsequent multicenter study by an international consortium of clinical facilities that had collected both brain FDG-PET and histopathologic data for patients undergoing evaluation for dementia,31 AD was identified using PET with sensitivity and specificity of 94% and 73%, respectively. This latter study, which included more than 3 times as many patients as the 4 previous series combined, included a stratified examination of the subset of patients with documented early or mild disease for whom performance of PET with

**Figure 3** Changes in cortical metabolism typical for various degrees of impairment, from normal to late AD. FDG-PET images of transaxial planes from 5 patients, shown at comparable axial levels. First row, Normal pattern, provided for reference. Note how posterior cingulate cortex (arrow) normally has activity that is visibly higher than in average cortex. Second row, Patient with age-associated memory impairment not meeting the criteria for MCI. Arrow denotes patchiness in the inferior parietal cortex that is beginning to emerge, and activity in posterior cingulate cortex is also seen to be less robust than in normal subject. Third row, MCI, with clear hypometabolism of parietal, parietotemporal and posterior cingulate cortex. Fourth row, Early AD, demonstrating posterior-predominant cortical hypometabolism. Fifth row, Late AD, with bilateral precortical, parietal, temporal, and posterior cingulate cortical regions markedly hypometabolic, but with continued relative preservation of sensorimotor and visual cortex. At lower planes than shown here, basal ganglia, thalamus, cerebellum and brainstem would also be seen to be relatively preserved at all stages. (Color version of figure is available online.)
respect to sensitivity (95%), specificity (71%), and overall accuracy (89%) was nearly the same. The aforementioned values are in accordance with the ranges found in a broader review of the PET literature, which additionally included studies lacking neuropathologic confirmation of diagnoses,\(^8\) that found sensitivities ranging from 90% to 96% and specificities ranging from 67% to 97%, as well as in an assessment of the PET literature reported by the American Academy of Neurology,\(^8\) based on their review of published studies which demonstrated diagnostic accuracies of 86-100% for PET.

**Other Tracers**

Although clinical applications of PET in the evaluation of patients with conditions leading to cognitive impairment and dementia have largely focused on FDG, many other tracers have been used in this context. Although a detailed consideration of those tracers is beyond the scope of the present review, a few such examples will be mentioned here. To begin, the use of PET to study cerebrovascular disease, as with AD, also enjoys a long history, extending back to the time of the earliest dementia studies.\(^8\) These conditions overlap, in that uncontrolled chronic cerebrovascular disease can eventually lead to vascular dementia (Fig. 5). PET has been used to directly quantify several parameters pertinent to the status of the cerebrovascular system, including cerebral blood flow, cerebral blood volume, and cerebral rate of oxygen metabolism, using \(^{15}\)O-labeled water, carbon monoxide, and oxygen gas. This has allowed estimation of further relevant parameters through calculations based on the values derived from those measurements, including cerebrovascular mean transit time, cerebral perfusion pressure, oxygen extraction fraction, and stoichiometry of oxygen and glucose utilization. Each of these has been reported to change, in different ways, under circumstances stemming from the pathophysiological events that occur during cerebrovascular compromise, and the evolution of stroke as well as its aftermath.

For example, oxygen extraction fraction data, calculated from PET measures of regional oxygen metabolism and blood flow using \(^{15}\)O-oxygen gas and \([H_2]^{15}\)O-water, respectively, were demonstrated to predict stroke risk in patients having a history of stroke or transient ischemic attack in the distribution of an occluded carotid artery.\(^8\) Patients with increased oxygen extraction fractions (operationally defined by asymmetry, through reference to a control group, approximately corresponding to exceeding the contralateral region by more than 8%), after adjusting for age, had a 6-fold greater risk of suffering a stroke (all but one occurring ipsilateral to the side

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**Figure 4** Dementia with Lewy bodies. FDG-PET pattern is typically posterior-predominant, as in AD, but without preferential sparing of occipital cortex.

**Figure 5** Vascular dementia. Patient was diagnosed both clinically and by structural imaging with vascular dementia. Arrows on this scan indicate hypometabolism of the right parietal cortex (left), right prefrontal cortex, basal ganglia and thalamus (middle), and right temporal cortex (right). The hypometabolism of the left cerebellum (right) is characteristic of cross-cerebellar diaschisis, caused by diminished afferent input from contralateral cortex.
with higher oxygen extraction fraction) than controls. It was independently found that increased hemispheric oxygen extraction (defined by a reference value derived from the mean and variance of the extraction fraction of a control group, corresponding to exceeding 53.3%) was associated with a 7-fold greater risk of suffering a stroke. (A less significant difference was found between groups categorized according to asymmetry of oxygen extraction fraction.)

In the setting of acute stroke, restoration of cerebral blood flow demonstrated with $^{15}$O-water, in patients scanned before and after thrombolytic therapy with tissue plasminogen activator (tPA), predicted clinical improvement assessed 3 weeks later. In PET studies performed within 5 to 18 hours of onset of middle cerebral artery stroke, the extent of abnormally low cerebral blood flow or oxygen metabolism (but not blood volume, perfusion pressure, or oxygen extraction fraction) correlated with final infarct size and long-term clinical outcome. Beyond investigation of these relatively well-studied hemodynamic parameters measurable with PET, studies with newer tracers (eg, those recognizing activated microglial cells or benzodiazepine receptors) are also showing promise of contributing clinically meaningful information to the assessment of patients with cerebrovascular disease. Recent studies have also turned to comparing the aforementioned hemodynamic parameters measured with PET alone to parameters obtained with use of acetazolamide or obtained with perfusion-weighted MRI. It remains to be seen what impact any of these methods to assess various aspects of cerebral status will have on routine clinical management of patients with cerebrovascular disease.

Much of the attention of the past few years in dementia imaging has been focused on imaging of amyloid plaques and neurofibrillary tangles with PET. Just as changes in glucose metabolism occur early in the development of AD pathology, evidence from postmortem studies indicates that the accumulation of senile plaques and neurofibrillary tangles, the neuropathological hallmarks of AD, occurs over the decades before clinical AD diagnosis. In an effort to visualize these neuropathological lesions in vivo, several groups have independently developed PET tracers with which to label these pathologies. Best studied in humans are the thioflavin T derivative 11C-PIB, which recognizes beta-amyloid plaques, and 18F-FDDNP, which labels plaques and tangles. Both have been used in patients with AD and with MCI. These types of compounds may prove particularly useful in drug development, in particular for those drugs that are aimed at diminishing the burden of insoluble plaques or neurofibrillary tangles in the brain.

Many other tracers have been aimed at neurotransmitter systems affected by Alzheimer’s and other neurodegenerative diseases, including probes for cholinergic receptors and cholinesterase enzymes, dopaminergic transporters, and serotonin receptors. For example, reduced density of medial temporal serotonin receptors can serve as an index of hippocampal pyramidal neuronal loss, and 18F-MPPF is a selective imaging probe for 5-HT1A receptors in hippocampus and other brain areas. This permits quantification of 5-HT1A receptors in the human brain with PET, in a manner that distinguishes patients with AD from normal controls. Although many of these tracers have been useful in uniquely enabling examination of molecular processes associated with dementia in living brain tissue, it again remains to be seen what role they may play clinically.

**Future Directions**

It is evident that the biologic changes that occur with AD take place decades before the onset of clinical symptoms. The use of more powerful diagnostic approaches allows the earlier detection of initial changes in a target population long before a diagnosis of AD can be made. Currently, the availability of tools for identifying the early changes of AD is outpacing the available therapeutic options. With advances in potential disease-modifying treatments, the benefits of early detection will become of greater importance, allowing therapy to be initiated as early as possible.

Much attention is currently focused upon methods intended to prevent AD or to modify its course. Given the low incidence and slow progression of normal elderly to AD (1-3%/yr), using traditional clinical endpoints to test the efficacy of such treatments would require very large samples, long follow-up, and great expense to follow cognitively normal persons treated with a candidate primary prevention therapy to assess impact on development of AD. FDG-PET imaging may be used to expedite the process by detecting brain abnormalities in individuals who might be at heightened risk for AD but who have not yet developed symptoms. Some studies used statistical power analyses to test the feasibility of using brain glucose metabolism as outcome measure in long-term treatment studies of AD in comparison with cognitive test scores. For example, it was estimated that to detect a 33% treatment response with 80% power in a typical 1-year, double-blind, placebo-controlled treatment study, a cognitive study using the Mini Mental State Examination would require 224 AD patients per group, compared with 36 patients per group would be needed for an FDG-PET study, due to the relatively small variance in regional brain metabolism measures from session to session, compared with the variance in neuropsychologic test performance.

Finally, PET may be used to examine cerebral responses to nonpharmacological preventative strategies. As an example of this approach, effects of a lifestyle intervention, consisting of healthy diet, exercise, and cognitive stimulation, on brain function and other physiological indices related to cardiac and brain health were measured. Healthy volunteers were randomized to the active intervention or a control activity. The intervention group showed significant changes in frontal lobe activity on PET compared with the control group that suggested increased efficiency in regional brain function. Larger clinical trials are underway to further assess the effects of such interventions.

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