The Differential Diagnosis of Alzheimer’s Disease: Conceptual and Methodological Issues

Mary C. Tierney, David W. Reid, Maria L. Zorzitto, W. Gary Snow, Rory H. Fisher, Irene Campbell-Taylor, and Anthony Lewis

ABSTRACT: The study of Alzheimer’s disease is hampered by insufficient knowledge of its cause. It can best be described as a syndrome whose clinical and pathological features, and their associations over time, need to be more carefully examined. Issues which impede our understanding of this syndrome include the lack of: (a) a singular “gold standard” for its identification; (b) longitudinal studies with appropriate comparison groups and neuropathological follow-up; and (c) standardized multifaceted clinical assessment procedures. Our awareness of the significance of these issues has led us to undertake a large-scale prospective, longitudinal investigation of 399 dementing and normal individuals at Sunnybrook Medical Centre. As a result of problems identified, it is proposed that research efforts across various Canadian centres be coordinated to best utilize available resources and expertise.

This presentation will deal with the contribution of research to the understanding of Alzheimer’s disease. Factors limiting progress and potential solutions to advance knowledge will be outlined. A review of the literature indicates that researchers usually operate on the assumption that Alzheimer’s disease is identifiable, either clinically or neuropathologically, and represents a distinct phenomenon. It is our contention that much of the research in Alzheimer’s disease is directed at attempts to explain a syndrome we do not fully understand. Yet, theoretical models are currently being developed to explain specific aspects of this poorly defined syndrome. While theory is important at a stage of research where the disease is well identified, premature testing of theory may lead to unproductive tangents.

One of the major limitations in research on Alzheimer’s Disease is that it has been largely compartmentalized within research specialties, such as neurology, neurochemistry, neuropsychology, epidemiology, and radiology. The merging of the substantive findings across research specialties will pose a major challenge. This can be met ideally by either collecting the data on the same subjects, or by using a common rigorous set of criteria across studies. While the former is unrealistic, the latter has already been suggested by the NINCDS and ADRDA Work Group but further refinement of these criteria is needed. Presently, these criteria do not describe specific markers for the disease, but rather they are mainly exclusionary and lack adequate standardization. As was recommended by this Work Group, these tentative criteria require longitudinal study and need to be evaluated in the context of pathological follow-up.

Unfortunately, research is rarely directed at the issue of reliability of identification at either the clinical or neuropathological levels. Assumptions of reliable identification are typically made on the basis of “neuropathological verification”. If Alzheimer’s syndrome can only be recognized at the neuropathological level, then this begs the question as to what is being identified, particularly because the diagnosis of a dementia remains a clinical one. If the neuropathologist must rely on clinical information to confirm the diagnosis of a dementia, the circularity becomes obvious. A clinical diagnosis requires neuro-
pathological confirmation and a neuropathological diagnosis requires clinical confirmation. This problem arises because, as with the clinical criteria, there are no specific neuropathological markers for Alzheimer’s syndrome. The brain pathologies considered characteristic of Alzheimer’s disease may also appear in the brains of nondemented elderly individuals. Thus, there is a demonstrated need to direct research efforts at a better understanding of the syndrome and ultimately to a better understanding of the brains of nondemented elderly individuals. Thus, there is a demonstration need to direct research efforts at a better understanding of the syndrome and ultimately to a better understanding of the brains of nondemented elderly individuals.

The next section of this paper will describe research strategies which, if employed, should lead to a more complete description of the syndrome and ultimately to a better understanding of it.

Sample size Because Alzheimer’s is a progressively deteriorating condition and may result in individual variation in expression, 3,4 large samples of individuals who meet the existing diagnostic criteria must be followed. This is necessary in order to determine whether there are common clinical features consistent across these individuals yet distinct from both normal aging and other dementing processes. Unfortunately, sample sizes have typically been too small to identify the differential patterns unique to Alzheimer’s disease.

Comparison groups In order to establish the distinguishing characteristics of Alzheimer’s disease at both the neuropathological and clinical levels, comparisons must be made among individuals with diagnoses of Alzheimer’s disease, diagnoses of dementias of other etiologies, as well as normals. The majority of studies examining the distinguishing characteristics of Alzheimer’s disease have focused on differences between the latter and normals. 5-7 Unfortunately, with few exceptions, those studies comparing Alzheimer’s with other dementias have only used small samples. 8-10 Because of these shortcomings, the value of the various measures in differentiating between Alzheimer’s disease and other forms of dementia remains to be established. The latter distinction is crucial if we are to progress to the stage of developing specific markers for the disease.

Another important rationale for the inclusion of individuals with dementias of different etiologies in the study sample is that it permits a comparison of the neuropathological features of these different groups. An illustration of the importance of such comparisons may be found in the research investigating subcortical dementias. Researchers examining the brains of individuals with Parkinsonian Lewy bodies have reported the presence of pathological changes characteristic of Alzheimer’s in many of these brains. 11,12 Speculations regarding a common etiology may well hinder our progress in the understanding of this syndrome.

The next section of this paper will describe research strategies which, if employed, should lead to a more complete description of the syndrome and ultimately to a better understanding of it.

Range of measurement Just as it is important to include appropriate comparison groups, it is also essential that individuals within one study are assessed with a broad range of clinical, laboratory and neuropathological measures. This will permit the examination of the interrelationships between scores on these measures and the determination of which measures identify differences among the dementias. As was mentioned earlier, the compartmentalization of research specialties at this stage may well hinder our progress in the understanding of the disease.

Sunnybrook Medical Centre Dementia Study

To illustrate the recommended research strategies, the following brief description of the Sunnybrook Dementia Study is provided. While this study does not incorporate all aspects, it represents a significant portion of the ideal approach to the study of Alzheimer’s disease. This study is a prospective, longitudinal investigation of 399 individuals which commenced in 1982. After careful screening and application of the recommended diagnostic criteria, 14 the following groups were included in the study: (1) Neurologically Normal (N = 120); (2) Pure Alzheimer (N = 91); (3) Other Dementias (N = 67); (4) Mixed Alzheimer (N = 64); (5) Non-Demented (not neurologically normal) (N = 57).

Each of the 399 individuals have undergone the following clinical assessments at 12 month intervals: (a) neurological assessment of disorientation reflexes; deep tendon reflexes; muscle tone and movements; bradykinesia; power; and extra ocular movements; (b) neuropsychological assessment of the following functions: memory, intelligence, language, tactile, motor, auditory, psychomotor, and visuoperception; (c) language assessment including picture description, sentence disambiguation, and story retelling; and (d) behavioural assessment of mental disability, physical disability, socially irritating behaviour, and disengagement.

In addition, the morphometric assessments of autopsied brains include the evaluation of over 80 representative sections from frontal, temporal and parietal lobes, and including hippocampus, corpus callosum, fornix, midbrain, medulla and cerebellum.

The data generated from the extensive measurement gathered from this large sample have lent themselves to detailed multivariate analyses. To meet the research strategies recommended in this paper, a research project such as this would also benefit from involvement of other health science specialties, such as epidemiology, neurochemistry and radiology. The coordination of these differing specialties would necessitate the use of the same diagnostic criteria, making it possible to corroborate the respective findings. To make this approach feasible it would also be necessary to coordinate research efforts across various Canadian centres as well as specialties, thus permitting the best utilization of resources and providing a national basis for substantial advancement of knowledge of Alzheimer’s disease.
ACKNOWLEDGEMENTS

This research was supported by the Ontario Ministry of Health, the Gerontology Research Council of Ontario, the Canadian Geriatric Research Society, and the Sunnybrook Medical Centre Research Fund. We are grateful to G. Nadon, MD, who provided diagnostic assessments, and the nursing staff of the Department of Extended Care, Sunnybrook Medical Centre.

REFERENCES