HUNTINGTON'S CHOREA IN RELATION TO HEREDITY AND EUGENICS

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It is now generally conceded that predisposition plays an important rôle in mental disorder but the views as to the nature of its part are very diverse and hazy. Some investigators conclude that there is an inheritance of a particular type of mental trouble, while others conceive that only a general psychopathic constitution is inherited. Again a difference of view maintains as to the nature of the recognized mental 'diseases'—by some these are regarded as distinct 'entities,' manifestations of a single, indivisible, disturbing factor. Others conceive of them as syndromes or chance associations of symptoms. Even those who hold the latter view would perhaps except as the one clear case of a neuropathic entity the condition known as 'Huntington's Chorea.' This chorea is defined by the following traits: (1) persistent tremors of the head, appendages and trunk; (2) the onset of such tremors in middle or late life; (3) the progressive nature of the tremors; and (4) progressive mental deterioration. These characters are frequently found together; is their association a necessary one?

A study of four family complexes in eastern Long Island, south-western Connecticut, south-central Connecticut and eastern Massachusetts which show nearly a thousand cases of Huntington's chorea yields the
remarkable result that practically all can be traced back to some half-dozen individuals, including three (probable) brothers who migrated to America during the 17th century. But, already, numerous ‘biotypes’ having specific and differential hereditary behavior have appeared. Thus there is a biotype in which the tremors are absent but mental deterioration present; a biotype in which the tremors are not accompanied by mental deterioration; a biotype in which the chorea does not progress; and a biotype in which the onset of the choreic movements is in early life. In general, the symptomatology of chronic chorea is dissimilar in different strains of families. The age of onset, the degree of muscular involvement, the extent of mental deterioration all show family differences and enable us to recognize various species, or biotypes, of the disease. These biotypes are less striking than they would be were it not for the extensive hybridization that is taking place between biotypes in random human matings.

The method of inheritance of some of the elements of Huntington’s chorea has been worked out. In general, the choreic movements never skip a generation and in other respects show themselves clearly to be a dominant trait. The mental disorder is usually of the hyperkinetic or manic type and this also shows itself as a dominant. The age of onset apparently tends to diminish in successive generations—‘law of anticipation’—but this is partly, if not wholly, illusory and is due to the fact that in comparing the age of onset in grandfathers with that in grandchildren we are not comparing on the same basis, for the grandparents are a selected lot (selected on the basis of late onset—at least late enough for them to become parents), while grandchildren include those in whom the onset is so early in life that they will never marry. If instead of comparing the average age of onset in successive generations, one compares the age of onset in a number of choreic parents, their parents and their grandparents, then the evidence for anticipation vanishes. Eight such series give for average age of onset of the propositus 35.5 years, parent 38.8, grandparents 36.9. In this series we can see no evidence of anticipation.

Among the 3000 odd relatives of the 962 choreics studied many nervous traits are recorded. Thus epilepsy is recorded 39 times, infantile convulsions 19 times, meningial inflammations and brain fever 51 times, hydrocephaly 41 times, feeblemindedness 73 times, Sydenham’s chorea 11 times, and tics 9 times, mostly in one small family. This incidence, which would seem high for an unselected population, suggests that chorea occurs in families characterized by a general liability to nervous and mental troubles.
Though it can be shown that the 962 cases of chorea originated from 6 or 7 ancestors and that the tendency has been handed down almost without a break through the generations and that for generations there have been individuals who recognized the hereditary nature of the disease and were influenced in marriage accordingly; nevertheless, there is no clear evidence that persons belonging to the choreic lines voluntarily abstain to any marked degree from, or are selected against, in marriage.