

CHAPTER 12

Natural History of Cerebral Palsy

ABSTRACT: The natural course of CP has changed greatly during the past 50 years. If appropriate health care is available, affected children without significant comorbidities have actuarial survival approaching that of the general population.

An understanding of the natural history of CP is essential for a proper prognosis and provision of support services. The natural history of untreated CP is one of progressive deterioration. It has been shown that without intervention, detrimental changes in gait and function will occur over time span as short as 1.5 years. The impact of skeletal growth during childhood can compound the primary neurological problem if muscles fail to lengthen in proportion to their adjacent long bones; and spastic muscles have been shown to grow more slowly than normal. Therefore, although CP is by definition a static neurological lesion and a disorder of motor function, the phenotype has also been labelled a 'progressive neuromuscular deformity'. During adolescent growth spurt, many patients gain substantial weight, their gait worsens and become wheelchair bound. Important items include prognosis for life expectancy, walking, and hand function. The symptoms range from a mild gait abnormality to profound total body involvement. The standardized mortality ratios for CP in all ages, is significantly raised. There is a sharp fall in SMR after 14 years though mortality remains higher from ischemic heart disease and cancer than in the general population.

KEYWORDS Predicting the ability to walk, Walking beyond the second decade of life, Hand function: use it or lose it, Life expectancy, Quality of life and costs of caring, Medicolegal aspects

An understanding of the natural history of CP is essential for a proper prognosis and provision of support services. The natural history of untreated CP is one of progressive deterioration. It has been shown that without intervention, detrimental changes in gait and function will occur over time span as short as 1.5 years. During adolescent growth spurt, many patients gain substantial weight, their gait worsens and become wheel-chair bound. Important items include prognosis for life expectancy, walking, and hand function.

The impact of skeletal growth during childhood can compound the primary neurological problem

if muscles fail to lengthen in proportion to their adjacent long bones; and spastic muscles have been shown to grow more slowly than normal. Therefore, although CP is by definition a static neurological lesion and a disorder of motor function, the phenotype has also been labeled a 'progressive neuromuscular deformity' (Graham 2002). The symptoms range from a mild gait abnormality to profound total body involvement.

Survival

Children with total body involvement and bulbar difficulties, necessitating alternative feeding, are

Natural History Of Cerebral Palsy

DP Hallahan



Natural History Of Cerebral Palsy:

Whispering the Techniques of Language: An Emotional Journey through **Natural History Of Cerebral Palsy**

In a digitally-driven world where screens reign supreme and immediate connection drowns out the subtleties of language, the profound techniques and psychological nuances hidden within phrases frequently get unheard. However, situated within the pages of **Natural History Of Cerebral Palsy** a interesting fictional treasure blinking with raw thoughts, lies a fantastic quest waiting to be undertaken. Written by an experienced wordsmith, that charming opus encourages visitors on an introspective trip, softly unraveling the veiled truths and profound influence resonating within the very cloth of each word. Within the psychological depths with this emotional review, we will embark upon a genuine exploration of the book is core subjects, dissect their fascinating writing fashion, and succumb to the strong resonance it evokes heavy within the recesses of readers hearts.

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