Hŏnpŏp chaep'an yŏn'gu

Ch'ŏrhak kwa Hyŏnsilsa - MUSCLE AND NERVE DISORDERS IN CHILDREN

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Hŏnpŏp chaep'an chŏlch'a ŭi kaesŏn ŭl wihan ippŏmnonjŏk yŏn'gu. (1993 edition)

As soon as scoliosis is recognized, institute treatment by linearly posturing the spine on pads when supine and, if respiratory capacity permits, by a thoracolumbosacral orthosis with abdominal relief.

Report: PMP22 Variant Causes Peripheral Neuropathy in CMT Patients

The lack of protective sensation allows unrestricted repetitive trauma, and large weight-bearing joints are most frequently and severely involved. The CPK level is markedly elevated, with levels similar to those seen in Duchenne muscular dystrophy, and the results of muscle biopsy resemble those in Duchenne muscular dystrophy. A nerve biopsy may aid in the diagnosis of a peripheral neuropathy but is rarely required.

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The hamstrings are released distally, and the tendo Achilles is lengthened by a percutaneous method Fig.

MUSCLE AND NERVE DISORDERS IN CHILDREN

However, the prognostic value of p53 and Ki67 in fiberoptic bronchial biopsies FBB has not been fully investigated. Ambulatory patients have only mild contractures, which seldom interfere with walking. HNPCC is a relatively rare disease, which makes screening the entire populace burdensome and ineffective.

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The investigators compared the predictive value of the genotype score to that of family history alone or of physiological risk factors. Ten days later, the casts are removed, long-leg orthoses are measured, and casts are reapplied until the orthoses are fabricated. A cavus foot has a pathologic elevation of the longitudinal arch of less than 150° on the lateral radiograph at the intersection of the axis of the first metatarsal and the calcaneus 169.

Report: PMP22 Variant Causes Peripheral Neuropathy in CMT Patients

The gene abnormality has been localized to 4Q35, but no gene product has been identified 114, 365.

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