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Altri autori (Persone)	MerrittH. Houston <1902-1979.> (Hiram Houston)
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Soggetti	Nervous system - Diseases Neurology Electronic books.
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Nota di bibliografia	Includes bibliographical references and index.
Nota di contenuto	Signs and symptoms in neurologic diagnosis: approach to the patient -- Delirium and confusion -- Memory loss, behavioral changes and dementia -- Aphasia, apraxia, and agnosia -- Syncope, seizures and their mimics -- Coma -- Headache -- Diagnosis of pain and paresthesias -- Dizziness, vertigo, and hearing loss -- Impaired vision -- Involuntary movements -- Syndromes caused by weak muscles -- Gait disorders -- CT and MRI -- Electroencephalography and evoked potentials -- Electromyography, nerve conduction studies, and magnetic stimulation -- Autonomic testing -- Neurovascular imaging -- Endovascular neuroradiology -- Lumbar puncture and cerebrospinal fluid examination -- Muscle and nerve biopsy -- Neuropsychological evaluation -- DNA diagnosis -- Bacterial infections -- Focal infections -- Viral infections and postviral syndromes -- Human immunodeficiency virus (HIV) and acquired immunodeficiency syndrome (AIDS) -- Fungal infections -- Neurosarcoidosis -- Neurosyphilis -- Leptospirosis -- Lyme disease -- Parasitic infections -- Bacterial toxins -- Prion diseases -- Whipple disease -- Pathogenesis, classification, and epidemiology of cerebrovascular disease -- Examination of the patient with cerebrovascular disease -- Transient ischemic attack -- Cerebral infarction -- Intracerebral

hemorrhage -- Genetics of stroke -- Other cerebrovascular syndromes -- Differential diagnosis of stroke -- Stroke in children -- Treatment and prevention of stroke -- Subarachnoid hemorrhage -- Cerebral venous and sinus thrombosis -- Vascular disease of the spinal cord -- Vasculitis -- Susac syndrome -- Vascular tumors and malformations -- Hydrocephalus -- Normal pressure hydrocephalus (NPH) -- Brain edema and disorders of intracranial pressure -- Superficial siderosis and intracerebral hypotension -- Hyperosmolar syndromes -- General considerations -- Tumors of the skull and cranial nerves -- Tumors of the meninges -- Gliomas -- Lymphomas -- Pineal region tumors -- Tumors of the pituitary gland -- Congenital and childhood tumors -- Metastatic tumors -- Spinal tumors -- Paraneoplastic syndromes -- Complications of cancer chemotherapy --
Head injury -- Spine injury -- Cranial and peripheral nerve lesions -- Complex regional pain syndrome -- Radiation injury -- Electrical and lightning injury -- Decompression sickness -- Intervertebral dics and radiculopathy -- Cervical spondylotic myelopathy -- Thoracic outlet syndrome -- Hereditary and acquired spastic paraplegia -- Syringomyelia -- Neonatal neurology -- Floppy infant syndrome -- Disorders of motor and mental development -- Autism spectrum disorders -- Laurence-Moon-Biedl syndrome -- Cerebral and spinal malformations -- Chromosomal diseases -- Marcus Gunn -- Mobius syndrome -- Disorders of amino acid metabolism -- Disorders of purine and pyrimidine metabolism -- Lysosomal and other storage diseases -- Disorders of carbohydrate metabolism -- Glucose transporter type 1 deficiency syndrome -- Disorders of DNA maintenance, transcription, and translation -- Hyperammonemia -- Peroxisomal diseases: adrenoleukodystrophy, zellweger syndrome, and refsum disease -- Organic acidurias -- Disorders of metal metabolism -- Acute intermittent porphyria -- Neurologic syndromes with acanthocytes -- Cerebral degenerations of childhood -- Diffuse sclerosis and vanishing white matter disease -- Mitochondrial encephalomyopathies: diseases of mitochondrial DNA -- Leber hereditary optic neuropathy -- Mitochondrial diseases with mutations of nuclear DNA -- Neurofibromatosis -- Tuberous sclerosis complex -- Encephalotrigeminal angiomatosis -- Incontinentia pigmenti -- General considerations -- Alzheimer disease -- Frontotemporal dementia -- Lewy body dementias -- Huntington disease -- Choreas -- Myoclonus -- Gilles de la tourette syndrome -- Dystonia -- Essential tremor -- Parkinson disease -- Parkinson-plus syndromes -- Paroxysmal dyskinesias -- Tradive dyskinesia and other neuroleptic-induced syndromes -- Autosomal recessive ataxias -- Autosomal dominant ataxias -- Amyotrophic lateral sclerosis, progressive muscular atrophy, and primary lateral sclerosis -- Kennedy disease -- Spinal muscular atrophies of childhood -- Monomelic muscular atrophy -- General considerations -- The inherited peripheral neuropathies -- Acquired neuropathies -- Neuropathic pain -- Myasthenia gravis -- Lambert-Eaton syndrome -- Botulism and antibiotic-induced neuromuscular disorders -- Critical illness myopathy and neuropathy -- Identifying disorders of the motor unit --
Progressive muscular dystrophies -- Familial periodic paralysis -- Congenital disorders of muslce -- Myoglobinuria -- Muscle cramps and stiffness -- Dermatomyositis -- Polymyositis, inclusion body myositis, and related myopathies -- Myositis ossificans -- Multiple sclerosis -- Neuromyelitis optica -- Marchiafava-Bignami disease -- Central pontine myelinolysis -- Epilepsy -- Febrile seizures -- Primary and secondary headaches -- Transient global amnesia -- Meniere syndrome -- Sleep disorders -- Neurogenic orthostatic hypotension,

autonomic failure, and autonomic neuropathy -- Familial dysautonomia -- Endocrine diseases -- Hematologic and related diseases -- Hepatic disease -- Cerebral complications of cardiac surgery -- Bone disease -- Renal disease -- Respiratory support for neurologic diseases -- Nutritional disorders: malnutrition, malabsorption, and B and other vitamin deficiency -- Hypertrophic pachymeningitis -- Neurologic disease during pregnancy -- Hashimoto encephalopathy -- Schizophrenia -- Mood disorders -- Anxiety disorders -- Somatoform disorders -- Alcoholism -- Drug dependence -- Iatrogenic disease -- Occupational and environmental neurotoxicology -- HIV, fetal alcohol and drug effects, and the battered child -- Falls in the elderly -- Neurologic rehabilitation -- End-of-life issues in neurology.

Sommario/riassunto

The thoroughly updated Twelfth Edition of this classic retains the organization, practicality, and readability that makes Merritt's Neurology one of the most popular texts among neurologists, primary care physicians, and residents reviewing for psychiatry or neurology boards. In 183 short chapters, the book provides the essentials clinicians need on symptoms/signs, diagnostic tests, and neurologic disorders of all etiologies. For this edition, Timothy A. Pedley, MD joins Dr. Rowland as co-editor. Coverage includes separate chapters on autism, autosomal recessive ataxias, and autosomal dominant ataxias, and new chapters on endovascular neuroradiology, parkinsonian syndromes, Lewy body dementias, frontotemporal dementia, vanishing white matter, vasculitis, normal pressure hydrocephalus, neuromyelitis optica, Kennedy disease, spinal muscular atrophy, complex regional pain syndrome, and Hashimoto encephalopathy. A companion Website will offer the fully searchable text and an image bank.