

Enfermedad De Huntington Ppt

Johanna Knowles

Manual de Diagnósticos de Enfermería Betty J. Ackley, Gail B. Ladwig, 2006-12 Incluye la información más actual sobre los diagnósticos de enfermería 2005-2006 aprobados por la NANDA. También se incluyen las taxonomías de NIC y NOC, además de práctica basada en la evidencia, asistencia domiciliar y enseñanza a pacientes y familias. El método en tres pasos de Ackley y Ladwig enseña a relacionar los diagnósticos de enfermería con la información sobre los pacientes a partir de los resultados de la evaluación, los diagnósticos médicos, quirúrgicos o psiquiátricos y el plan de tratamiento actual para ayudar a priorizar e individualizar el plan de cuidados. Las tres secciones del libro formulan un diagnóstico de enfermería y crean un plan de cuidados utilizando los resultados importantes de la evaluación del paciente (sección I); proporcionan una lista de todos los diagnósticos de enfermería potenciales relacionados con el diagnóstico médico, los procedimientos diagnósticos, los estados clínicos y los síntomas del paciente (sección II), y ofrecen características definidas, factores de riesgo, intervenciones de la NIC y resultados de la NOC, y proporcionan justificaciones basadas en la evidencia para todas las intervenciones (sección III).

Rang y Dale. Farmacología James M. Ritter, Rod J. Flower, Graeme Henderson, Yoon Kong Loke, David MacEwan, Humphrey P. Rang, 2020-02-14 Nueva edición del texto de referencia en farmacología que, con una trayectoria de más de 25 años se convierte en la obra de primera elección para el estudiante de Medicina, por su carácter didáctico y abordaje visual. La nueva edición se organiza de la misma forma que la ed. vigente (6 secciones) a través de las cuales se explican los mecanismos de actuación de los fármacos. Presenta de una forma clara y exhaustiva los aspectos más relevantes de la farmacología, desde la comprensión de los receptores moleculares y las acciones farmacológicas, hasta las aplicaciones clínicas de grupos esenciales de fármacos. Se ha actualizado toda la obra, de forma que se recogen los últimos avances en la especialidad, incluidos los fármacos y las investigaciones más novedosas. Entre ellas cabe destacar una amplia revisión de los fármacos del SNC, los potenciadores cognitivos, los antiinfecciosos, los productos biológicos (biofármacos) y los fármacos extraterapéuticos. Acceso a un banco de preguntas de autoevaluación y a casos clínicos desarrollados.

All Within Susan E. Lawrence, 2012-07-01 Susan Lawrence continues her drive, building awareness as a full time advocate for families living with Huntington's disease. All Within is a compilation of personal stories from individuals within the HD community. Susan has turned to the families to openly share their stories and now moves forward with her fifth

addition of writings about HD, sharing personal messages behind the struggles of many. HD is a genetic disorder that affects people worldwide. There are those who suffer from the adult form of Huntington's disease, and the Juvenile form that prevents children from being able to reach adulthood and live full lives. There is currently no cure! It is a frustrating battle for us to help the world understand that it is what's inside that prevents us from finding peace. That means if we are not proactive advocates and continue sharing our experiences, everything is at risk of remaining All Within.

Genes & Generations Alison Gray,1995

Huntington's Disease Clay Farris Naff,2012-06-12 Huntington's Disease causes degeneration of brain cells, in motor control regions of the brain, as well as other areas. Symptoms get progressively worse, and include uncontrolled movements, abnormal body postures, and changes in emotion, behavior, judgment, and cognition. People with HD also develop impaired coordination, slurred speech, and difficulty feeding and swallowing. Provide your readers with essential information on HD. This book also serves as a historical survey, by providing information on the controversies surrounding its causes. Compelling first-person narratives by people coping with Huntington's Disease give readers a first-hand experience. Patients, family members, or caregivers explain the condition from their own experience. The symptoms, causes, treatments, and potential cures are explained in detail. Essential to anyone trying to learn about diseases and conditions, the alternative treatments are explored. Student researchers and readers will find this book easily accessible through its careful and conscientious editing and a thorough introduction to each essay.

Huntington's Disease ,1981

Huntington's Disease Randall McPartland,2015-12-15 Huntington's disease is an incurable and fatal degenerative condition that robs people of normal brain function when they should be in the prime of life. This book describes the condition and the genetic causes behind it, follows researchers on their path to scientific discovery, identifies people with the condition who have spoken up for the good of others, and tracks the latest treatments and research aimed at helping those living with it. Sidebars highlight medical breakthroughs and the people who made them.

Huntington's Disease Johanna Knowles,2006-07 Describes the history, symptoms, and diagnosis of Huntington's disease.

Report United States. Commission for the Control of Huntington's Disease and Its Consequences,1978

A Physician's Guide to the Management of Huntington's Disease Adam Rosenblatt,1999-01-01

Huntington's Disease National Institutes of Health (U.S.),1979

Huntington's Disease Thomas J. Visser,2010-01-01 Huntington's disease, chorea or disorder (HD) is an incurable neurodegenerative genetic disorder, which affects muscle co-ordination and some cognitive functions, typically becoming noticeable in middle age. It is the most common genetic cause of abnormal involuntary writhing movements called chorea and is much more common in people of Western European descent than in those from Asia or Africa. The disease is caused by

a dominant mutation on either of the two copies of a gene called Huntington. This book gathers and presents current research in the study of Huntington's Disease including the outlining effects of mutant htt in the nucleus and cytoplasm and the role of cell-cell interactions in Huntington's Disease pathology, as well as a review of the role of Huntington (HTT) interacting proteins.

Huntington's Disease Jurgen Lemiere,2004 This is a Ph.D. dissertation. Huntington's Disease (HD) is a dominantly inherited late onset neurodegenerative disorder, initially affecting the basal ganglia and especially the head of the caudate nucleus. Eventually, atrophy is more diffuse extending to the cerebral cortex and other brain nuclei. A triad of progressive motor, cognitive, and emotional symptoms characterizes adult onset HD. Contents include: General Introduction and Aims of the Study, Cognitive Changes in Patients with HD and Asymptomatic Carriers of the HD Mutation, Recent Memory and Recognition Memory in Huntington's Disease, Habitual Learning in Huntington's and Parkinson's Disease, Effect of Oral Creatine Supplementation on Cognition in HD, General Summary and Perspectives.

Huntington's Disease Kevin Guillory,2013

Living with Huntington's Disease Dennis H. Phillips,1982

Huntington's Disease S. B. Dunnett,Anne Rosser,2009-08-19

Huntington's Disease ,1989

Huntington's Disease Christina Hughes,2017 Huntington's disease (HD) is a progressive, fatal, dominant inherited neurodegenerative disorder caused by an unstable expansion of CAG repeats within the coding region of the IT15 gene that encodes for a protein called huntingtin (htt). The clinical hallmark of HD is a severe motor impairment, but cognitive and psychiatric symptoms are early clinical features in HD that often appear before the onset of motor signs. This book discusses the prevalence, pathogenesis and treatment of HD.

Huntington's Disease ,2019

Overview United States. Department of Health, Education, and Welfare. Commission on the Control of Huntington's Disease and its Consequences,1977

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