1. Record Nr. UNINA9910437617803321 Autore Morgan Kevin Titolo Genetic variants in Alzheimer's disease / / Kevin Morgan Pubbl/distr/stampa New York, : Springer, c2013 **ISBN** 1-4614-7309-8 Edizione [1st ed. 2013.] Descrizione fisica 1 online resource (256 p.) Disciplina 616.831042 Soggetti Alzheimer's disease - Genetic aspects Neurogenetics Lingua di pubblicazione Inglese **Formato** Materiale a stampa Livello bibliografico Monografia Description based upon print version of record. Note generali Nota di bibliografia Includes bibliographical references and indexes. The Genetics of Alzheimer's disease: Introduction and Perspective for Nota di contenuto the Future -- Apolipoprotein E -- Clusterin -- PICALM -- Complement Component (3b/4b) Receptor 1(CR1) -- Bridging Integrator 1 (BIN1) --ATP-binding cassette, sub-family A (ABC1), member 7 (ABCA7) --Membrane-spanning 4-domains subfamily A, MS4A cluster -- Sialic acid binding immunoglobulin-like lectin-3 (CD33) -- Erythropoietinproducing human hepatocellular carcinoma (EphA1) -- CD2-associated protein (CD2AP) -- Other Genes Implicated in Alzheimer's Disease --The Future Role of Biomarkers in Alzheimer's Disease Diagnostics --Index. Since 2009, a revolution has been witnessed in Alzheimer's Disease Sommario/riassunto genetics. New genetic links are being discovered at an unprecedented pace and our understanding of the molecular mechanisms of neurodegeneration have taken a quantum leap forward. This book provides a thorough description of the genes that have been implicated in the aetiology of late-onset Alzheimer's disease (LOAD) based on evidence of genetic association. These "AD susceptibility genes" are described both in their genomic and cellular context, as well as with respect to their known or suspected molecular functions. Although these genes are not sufficient to explain all of the genetic contributions

> to LOAD, they represent the best replicated set of genes to date. Undoubtedly the list will grow as more advanced genomic approaches

towards the identification of novel LOAD genes progresses.