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- PROTEIN MISFOLDING, AGGREGATION AND CONFORMATIONAL DISEASES VLADIMIR N. UVERSKY, ANTHONY FINK, 2006-06-13 RESEARCH INDICATES THAT MOST NEURODEGENERATIVE DISEASES, SYSTEMIC AMYLOIDOSES AND MANY OTHERS, ARISE FROM THE MISFOLDING AND AGGREGATION OF AN UNDERLYING PROTEIN. THIS IS THE FIRST BOOK TO DISCUSS SIGNIFICANT ACHIEVEMENTS IN PROTEIN STRUCTURE-FUNCTION RELATIONSHIPS IN BIOCHEMISTRY, MOLECULAR BIOLOGY AND MOLECULAR MEDICINE. THE AUTHORS SUMMARIZE RECENT PROGRESS IN THE UNDERSTANDING OF THE RELATIONSHIPS BETWEEN PROTEIN MISFOLDING, AGGREGATION AND DEVELOPMENT OF PROTEIN DEPOSITION DISORDERS.
- 2 ALZHEIMER'S DISEASE: INSIGHTS INTO LOW MOLECULAR WEIGHT AND CYTOTOXIC AGGREGATES FROM IN VITRO AND COMPUTER EXPERIMENTS PHILIPPE DERREUMAUX, 2012-12-31 ALZHEIMER'S DISEASE IS THE MOST COMMON FORM OF SENILE DEMENTIA, AFFECTING MORE THAN 24 MILLION PEOPLE WORLDWIDE. IT IS CHARACTERISED PATHOLOGICALLY BY ABNORMALLY HIGH LEVELS OF NEUROFIBRILLARY TANGLES RESULTING FROM THE ACCUMULATION OF TAU PROTEIN IN DEAD AND DYING NEURONS, AND BY ELEVATED NUMBERS OF SENILE PLAQUES IN THE CORTEX and hippocampus of the brain. The major component of senile plaques is a small protein of 39-43 amino acids called amyloid-b (Ab). Thus far, no treatment has been SHOWN TO SLOW THE PROGRESSION OF SPORADIC AND FAMILIAL ALZHEIMER'S DISEASE. A LARGE BODY OF EVIDENCE POINTS, HOWEVER, TO THE EARLY AB-FORMED OLIGOMERS AS THE PRIMARY TOXIC SPECIES IN ALZHEIMER'S DISEASE. A POWERFUL STRATEGY FOR DEVELOPING PHARMACEUTICAL TREATMENTS AGAINST ALZHEIMER'S IS TO ELUCIDATE THE PATHWAYS OF OLIGOMER FORMATION AND DETERMINE THE STRUCTURES OF THE TOXIC AGGREGATES. THIS BOOK PROVIDES A PANORAMIC VIEW ACROSS RECENT IN VITRO AND IN VIVO STUDIES ALONG WITH STATE-OF-THE-ART COMPUTER SIMULATIONS, DESIGNED TO INCREASE THE READERS' UNDERSTANDING OF AB OLIGOMERISATION AND FIBRIL FORMATION. AT THE SAME TIME, THE BOOK DELVES INTO THE PATHOGENESIS OF FAMILIAL AND SPORADIC ALZHEIMER'S DISEASE AT THE ATOMIC LEVEL OF DETAIL. WRITTEN BY LEADING AUTHORS IN THEIR RESPECTIVE FIELDS, THIS BOOK WILL BE VALUABLE TO ALL SCIENTISTS WORKING ON ALZHEIMER'S DISEASE. CONTENTS: CELLULAR AND IN VITRO ASPECTS OF AMYLOID-BETA (AB) AGGREGATION: AMYLOID HYPOTHESIS: MOLECULAR AND CELLULAR ASPECTS OF TOXICITY (RAKEZ KAYED AND CRISTIAN A LASAGNA-REEVES) MODELS OF WILD-TYPE AND DISEASE-CAUSING MUTANT B-AMYLOID FIBRILS. INSIGHTS FROM SOLID-STATE NUCLEAR MAGNETIC RESONANCE SPECTROSCOPY (STEPHEN C MEREDITH) BIOPHYSICAL CHARACTERIZATION OF AB ASSEMBLY (ERIC Y HAYDEN AND DAVID B TEPLOW) COORDINATION OF METAL IONS TO B-AMYLOID PEPTIDE: IMPACT ON ALZHEIMER'S DISEASE (PETER FALLER, GIOVANNI LA PENNA, CHRISTELLE HUREAU AND SARA FURLAN) AMYLOIDOGENESIS, NEUROGENESIS, LEARNING, AND MEMORY IN ALZHEIMER'S DISEASE: LESSONS FROM TRANSGENIC MOUSE MODELS (LAURE VERRET AND CLAIRE RAMPON) INHIBITOR DESIGN AGAINST CYTOTOXIC B-AMYLOID SPECIES (ANDREW J DOIG) BIOINFORMATICS AND COMPUTER SIMULATIONS OF AB AGGREGATION UNDER VARIOUS ENVIRONMENTAL CONDITIONS:KINETICS OF AMYLOID GROWTH (JESSICA NASICA-LABOUZE AND NORMAND MOUSSEAU)PROBING THE STABILITY OF FIBRIL AND TUBULAR SPECIES USING ALL-ATOM MOLECULAR DYNAMICS SIMULATIONS IN SOLUTION: INSIGHT INTO POLYMORPHISM (YIFAT MILLER, BUYONG MA AND RUTH NUSSINOV) MECHANISMS OF GROWTH OF B-AMYLOID FIBRILS AND BINDING OF NON-STEROIDAL ANTI-INFLAMMATORY DRUG LIGANDS (TAKAKO TAKEDA AND DMITRI KLIMOV) EXPLORING THE STRUCTURES OF B-AMYLOID OLIGOMERS IN AQUEOUS SOLUTION USING COARSE-GRAINED PROTEIN MODELS (YASSMINE CHEBARO AND PHILIPPE DERREUMAUX)PATHWAYS OF AMYLOID FIBRIL FORMATION USING A SIMPLIFIED PEPTIDE MODEL (RICCARDO PELLARIN) FROM DISORDERED AMYLOID B-PROTEINS TO SOLUBLE OLIGOMERS AND PROTOFIBRILS USING DISCRETE MOLECULAR DYNAMICS (MARK BETNEL, NIKOLAY V DOKHOLYAN AND BRIGITA URBANC) AB PROTEINS-LIPID MEMBRANE INTERACTION: COMPUTATIONAL SIMULATION STUDY (YUGUANG MU) MOLECULAR INSIGHTS INTO THE ASSEMBLY OF B-AMYLOID ON SURFACES AND CARBON NANOTUBES (GUANGHONG WEI, YIN LUO AND ZHAOMING FU) INHIBITING PEPTIDE AND PROTEIN SELF-AGGREGATION: WHAT CAN SIMULATIONS TELL US? (JOAN-EMMA SHEA AND GIORGIO COLOMBO) READERSHIP: GENERAL PUBLIC, PHD STUDENTS, POSTDOCS, AS WELL AS ALL SCIENTISTS WORKING IN THE FOLLOWING FIELDS: ALZHEIMER'S DISEASE, AMYLOIDS, NEUROSCIENCE, OLIGOMERS, PROTOFIBRILS, FIBRILS, TOXICITY, STRUCTURES, DYNAMICS, KINETICS, THERMODYNAMICS, ALL-ATOM AND COARSE GRAINED SIMULATIONS, BIOPHYSICS, MEMORY, LEARNING, TRANSGENIC MOUSE MODELS, METAL IONS, MEMBRANE, NANOPARTICLES, COMPUTATIONAL CHEMISTRY, DRUG DESIGN AND BIOINFORMATICS. KEYWORDS:AMYLOID; ALZHEIMER'S DISEASE; OLIGOMERS; FIBRILS; TOXICITY; STRUCTURES; DYNAMICS; THERMODYNAMICS; SIMULATIONS; BIOPHYSICS; MEMORY; TRANSGENIC MOUSE

KEYWORDS: AMYLOID; ALZHEIMER S DISEASE; OLIGOMERS; FIBRILS; I OXICITY; STRUCTURES; DYNAMICS; I HERMODYNAMICS; SIMULATIONS; BIOPHYSICS; MEMORY; I RANSGENIC MOUSE

MODELS; METAL IONS; DRUG DESIGNKEY FEATURES: WRITTEN BY PROMINENT AUTHORS IN THEIR FIELDS THE FIRST OVERVIEW AND PERSPECTIVE ON THE STATE-OF-THE-ART IN VITRO, IN SILICO

AND IN VIVO EXPERIMENTS AT THE HEART OF ALZHEIMER'S DISEASE

PRECISION MEDICINE IN NEURODEGENERATIVE DISORDERS, 2023-02-14 PRECISION MEDICINE IN NEURODEGENERATIVE DISORDERS, PART ONE, VOLUME 192 IN THE HANDBOOK OF CLINICAL NEUROLOGY DEALS WITH THE WHY IN THE APPROACH TO SLOW THE PROGRESSION OF ACCELERATED BRAIN AGING. THIS VOLUME IS INTENDED TO PROVIDE A SCHOLARLY BACKGROUND ON THE FRAMEWORK, BASIC SCIENCE AND CONCEPTUAL PITFALLS RELATED TO DISEASE-MODIFYING EFFORTS IN PARKINSON'S, ALZHEIMER'S AND OTHER NEURODEGENERATIVE DISORDERS. AMONG TOPICS COVERED ARE DIFFERENT MODELS OF PRECISION MEDICINE, THE LUMPING-VERSUS-SPLITTING TENSION IN BIOMARKER DEVELOPMENT AND THERAPEUTICS, AND THE RATIONALE FOR REPLACING THE CONVERGENCE OF THE PREVAILING AUTOPSY-BASED NOSOLOGY OF NEURODEGENERATIVE DISEASES WITH THE DIVERGENCE OF A SYSTEMS BIOLOGY APPROACH TO HUMAN DISEASES. SPECIFIC CHAPTERS ARE DEDICATED TO THE PROMISE OF GENETIC SUBTYPES AND THE LESSONS IN DISEASE MODIFICATION OFFERED BY THE FIELDS OF ONCOLOGY AND CYSTIC FIBROSIS THAT CAN BE ADAPTED TO THE FIELD OF NEURODEGENERATION. MATCHING A BIOLOGY-CORRECTING THERAPY WITH THOSE BIOLOGICALLY SUITABLE TO BENEFIT FROM SUCH THERAPY REPRESENTS THE VISION

AND MISSION OF PRECISION MEDICINE, THE HIGHEST LEVEL OF PERSONALIZED MEDICINE. SUMMARIZES THEORY AND RESEARCH ON PRECISION MEDICINE IN NEURODEGENERATIVE DISORDERS COVERS BASIC BIOLOGY, CLINICAL TRIALS AND THERAPEUTICS INCLUDES DISEASE MECHANISMS, GENETIC SUBTYPES, AND MORE

- EFICIENT, VERSATILE AND RELATIVELY CHEAP METHOD FOR DISSECTING THE MOLECULAR PATHWAYS THAT ARE THE BASIS OF LIFE, AS WELL AS FOR INVESTIGATING AND POTENTIALLY RECTIFYING FAULTS IN THESE PATHWAYS THAT RESULT IN DISEASE. THIS BOOK REVIEWS HOW CRISPR-Cas9 and other genome editing techniques are advancing our understanding of Development and function in the nervous system, uncovering the molecular causes of neurological disorders and providing tools for gene therapy.
- DERUG DISCOVERY APPROACHES FOR THE TREATMENT OF NEURODEGENERATIVE DISORDERS ADEBOYE ADEJARE, 2016-09-20 DRUG DISCOVERY APPROACHES FOR THE TREATMENT OF NEURODEGENERATIVE DISORDERS: ALZHEIMER'S DISEASE EXAMINES THE DRUG DISCOVERY PROCESS FOR NEURODEGENERATIVE DISEASES BY FOCUSING SPECIFICALLY ON ALZHEIMER'S DISEASE AND ILLUSTRATING THE PARADIGM NECESSARY TO ENSURE FUTURE RESEARCH AND TREATMENT SUCCESS. THE BOOK EXPLORES DIAGNOSIS, EPIDEMIOLOGY, DRUG DISCOVERY STRATEGIES, CURRENT THERAPEUTICS, AND MUCH MORE TO PROVIDE A HOLISTIC APPROACH TO THE DISCOVERY, DEVELOPMENT, AND TREATMENT OF ALZHEIMER'S DISEASE. THROUGH ITS COVERAGE OF THE LATEST RESEARCH IN TARGETED DRUG DESIGN, PRECLINICAL STUDIES, AND MOUSE AND RAT MODELS, THE BOOK IS A MUST-HAVE RESOURCE FOR ALL PHARMACEUTICAL SCIENTISTS, PHARMACOLOGISTS, NEUROSCIENTISTS, AND CLINICAL RESEARCHERS WORKING IN THIS AREA. IT ILLUSTRATES WHY THESE DRUGS TEND TO FAIL AT THE CLINICAL STAGE, AND EXAMINES ALZHEIMER'S DISEASE WITHIN THE OVERALL CONTEXT OF IMPROVING THE DRUG DISCOVERY PROCESS FOR THE TREATMENT OF OTHER NEURODEGENERATIVE DISORDERS. PROVIDES A COMPILATION OF CHEMICAL CONSIDERATIONS REQUIRED IN DRUG DISCOVERY FOR THE TREATMENT OF NEURODEGENERATIVE DISORDERS EXAMINES DIFFERENT CLASSES OF COMPOUNDS CURRENTLY BEING USED IN DISCOVERY AND DEVELOPMENT STAGES EXPLORES IN VITRO AND IN VIVO MODELS WITH RESPECT TO THEIR ABILITY TO TRANSLATE THESE MODELS TO HUMAN CONDITIONS DISTILLS THE MOST SIGNIFICANT INFORMATION ACROSS MULTIPLE AREAS OF ALZHEIMER'S DISEASE RESEARCH TO PROVIDE A SINGLE, COMPREHENSIVE, AND BALANCED RESOURCE
- ENERTICS, NEUROLOGY, BEHAVIOR, AND DIET IN PARKINSON'S DISEASE COLIN R MARTIN, VICTOR R PREEDY, 2020-08-13 GENETICS, NEUROLOGY, BEHAVIOR, AND DIET IN PARKINSON'S DISEASE: THE NEUROSCIENCE OF PARKINSON'S DISEASE, VOLUME 2 PROVIDES A SINGLE SOURCE OF MATERIAL COVERING DIFFERENT SCIENTIFIC DOMAINS OF NEUROPATHOLOGY UNDERLYING THIS CONDITION. THE BOOK COVERS A WIDE RANGE OF SUBJECTS AND UNRAVELS THE COMPLEX RELATIONSHIPS BETWEEN GENETICS, MOLECULAR BIOLOGY, PHARMACEUTICAL CHEMISTRY, NEUROBIOLOGY, IMAGING, ASSESSMENTS, AND TREATMENT REGIMENS. IT FILLS A MUCH-NEEDED GAP AS A ONE-STOP SYNOPSIS OF EVERYTHING TO DO WITH THE NEUROLOGY AND NEUROSCIENCE RELATED TO PARKINSON'S DISEASE—FROM CHEMICALS AND CELLS TO INDIVIDUALS. IT IS AN INVALUABLE RESOURCE FOR NEUROSCIENTISTS, NEUROLOGISTS, AND ANYONE IN THE FIELD. OFFERS THE MOST COMPREHENSIVE COVERAGE OF A BROAD RANGE OF TOPICS RELATED TO PARKINSON'S DISEASE SERVES AS A FOUNDATIONAL COLLECTION FOR NEUROSCIENTISTS AND NEUROLOGISTS ON THE BIOLOGY OF DISEASE AND BRAIN DYSFUNCTION CONTAINS IN EACH CHAPTER AN ABSTRACT, KEY FACTS, MINI DICTIONARY OF TERMS, AND SUMMARY POINTS TO AID IN UNDERSTANDING FEATURES PRECLINICAL AND CLINICAL STUDIES TO HELP RESEARCHERS MAP OUT KEY AREAS FOR RESEARCH AND FURTHER CLINICAL RECOMMENDATIONS SERVES AS A ONE-STOP SOURCE FOR EVERYTHING YOU NEED TO KNOW ABOUT PARKINSON'S DISEASE
- EMOSITRON EMISSION TOMOGRAPHY DALE L. BAILEY, DAVID W. TOWNSEND, PETER E. VALK, MICHAEL N. MAISEY, 2006-07-06 ESSENTIAL FOR STUDENTS, SCIENCE AND MEDICAL GRADUATES WHO WANT TO UNDERSTAND THE BASIC SCIENCE OF POSITRON EMISSION TOMOGRAPHY (PET), THIS BOOK DESCRIBES THE PHYSICS, CHEMISTRY, TECHNOLOGY AND OVERVIEW OF THE CLINICAL USES BEHIND THE SCIENCE OF PET AND THE IMAGING TECHNIQUES IT USES. IN RECENT YEARS, PET HAS MOVED FROM HIGH-END RESEARCH IMAGING TOOL USED BY THE HIGHLY SPECIALIZED TO AN ESSENTIAL COMPONENT OF CLINICAL EVALUATION IN THE CLINIC, ESPECIALLY IN CANCER MANAGEMENT. PREVIOUSLY BEING THE REALM OF SCIENTISTS, THIS BOOK EXPLAINS PET INSTRUMENTATION, RADIOCHEMISTRY, PET DATA ACQUISITION AND IMAGE FORMATION, INTEGRATION OF STRUCTURAL AND FUNCTIONAL IMAGES, RADIATION DOSIMETRY AND PROTECTION, AND APPLICATIONS IN DEDICATED AREAS SUCH AS DRUG DEVELOPMENT, ONCOLOGY, AND GENE EXPRESSION IMAGING. THE TECHNOLOGIST, THE SCIENCE, ENGINEERING OR CHEMISTRY GRADUATE SEEKING FURTHER DETAILED INFORMATION ABOUT PET, OR THE MEDICAL ADVANCED TRAINEE WISHING TO GAIN INSIGHT INTO THE BASIC SCIENCE OF PET WILL FIND THIS BOOK INVALUABLE. THIS BOOK IS PRIMARILY REPACKAGED CONTENT FROM THE BASIC SCIENCE SECTION OF THE 'BIG' VALK BOOK ON PET. IT CONTAINS NEW, COMPLETELY REVISED AND UNCHANGED CHAPTERS COVERING THE BASIC SCIENCES SECTION OF THE MAIN BOOK TOTAL 18 CHAPTERS: 2 NEW (CHAPTERS 1, 16) 8 COMPLETELY REVISED (CHAPTERS 4, 5, 8, 13, 14, 15, 17, 18) 3 MINOR CORRECTIONS (CHAPTERS 2, 6, 11) 5 UNCHANGED (CHAPTERS 3, 7, 9, 10, 12)
- FINEUROTHERAPEUTICS IN THE ERA OF TRANSLATIONAL MEDICINE. RICHARD A. SMITH, 2021
- PROTEIN MISFOLDING, AGGREGATION AND CONFORMATIONAL DISEASES VLADIMIR N. UVERSKY, ANTHONY FINK, 2007-05-26 THE SECOND VOLUME CONTINUES TO FILL THE GAP IN PROTEIN REVIEW AND PROTOCOL LITERATURE. IT DOES THIS WHILE SUMMARIZING RECENT ACHIEVEMENTS IN THE UNDERSTANDING OF THE RELATIONSHIPS BETWEEN PROTEIN MISFOLDINGS, AGGREGATION, AND DEVELOPMENT OF PROTEIN DEPOSITION DISORDERS. THE FOCUS OF PART B IS THE MOLECULAR BASIS OF DIFFERENTIAL DISORDERS.
- PAYNAPTIC PLASTICITY AND THE MECHANISM OF ALZHEIMER'S DISEASE DENNIS J. SELKOE, ANTOINE TRILLER, 2008-01-08 A BIOCHEMICAL HYPOTHESIS THAT ALZHEIMER'S DISEASE (AD)

IS A PROGRESSIVE CEREBRAL AMYLOIDOSIS CAUSED BY THE AGGREGATION OF THE AMYLOID B-PROTEIN (AB) - PRECEDED AND ENABLED THE DISCOVERY OF ETIOLOGIES. THIS VOLUME SERVES AS A RECORD FOCUSED ON BRINGING TOGETHER INVESTIGATORS AT THE FOREFRONT OF ELUCIDATING THE STRUCTURE AND FUNCTION OF HIPPOCAMPAL SYNAPSES WITH INVESTIGATORS FOCUSED ON UNDERSTANDING HOW EARLY ASSEMBLIES OF AB MAY COMPROMISE SOME OF THESE SYNAPSES.

- MAGING BRAIN DISEASES SERGE WEIS, MICHAEL SONNBERGER, ANDREAS DUNZINGER, MARTIN AICHHOLZER, RAIMUND KLEISER, PETER STRASSER, 2016-08-25 IMAGING BRAIN DISEASES ILLUSTRATES IN A UNIQUE WAY THE MOST COMMON DISEASES AFFECTING THE HUMAN NERVOUS SYSTEM USING DIFFERENT IMAGING MODALITIES DERIVED FROM RADIOLOGY, NUCLEAR MEDICINE, AND NEUROPATHOLOGY. THE FEATURES OF THE DISEASES ARE VISUALIZED ON COMPUTERIZED TOMOGRAPHY (CT)-SCANS, MAGNETIC RESONANCE IMAGING (MRI)-SCANS, NUCLEAR MEDICINE SCANS, SURGICAL INTRAOPERATIVE AS WELL AS GROSS-ANATOMY AND HISTOLOGY PREPARATIONS. FOR EACH DISEASE ENTITY, THE STRUCTURAL CHANGES ARE ILLUSTRATED IN A CORRELATIVE COMPARATIVE WAY BASED ON THE VARIOUS IMAGING TECHNIQUES. THE BRAIN DISEASES ARE PRESENTED IN A SYSTEMATIC WAY ALLOWING THE READER TO EASILY FIND THE TOPICS IN WHICH SHE OR HE IS PARTICULARLY INTERESTED. IN PART 1 OF THE BOOK, THE IMAGING TECHNIQUES ARE DESCRIBED IN A PRACTICAL, STRAIGHTFORWARD WAY. THE MORPHOLOGICAL BUILT-UP OF THE NORMAL HUMAN BRAIN AND ITS VASCULAR SUPPLY ARE PRESENTED IN PART 2. THE CHAPTERS OF THE SUBSEQUENT PARTS 3 TO 10 DEAL WITH THE FOLLOWING DISEASES INVOLVING THE NERVOUS SYSTEM INCLUDING: HEMODYNAMIC, VASCULAR, INFECTIOUS, NEURODEGENERATIVE, DEMYELINATION, EPILEPSY, TRAUMA AND INTOXICATION, AND TUMORS. THE AUTHORS INCITE THE CLINICIAN TO SEE THE CELL, THE TISSUE, THE ORGAN, THE DISORDER BY ENABLING HIM TO RECOGNIZE BRAIN LESIONS OR INTERPRETING HISTOLOGIC FINDINGS AND TO CORRELATE THIS KNOWLEDGE WITH MOLECULAR BIOLOGIC CONCEPTS. THUS, THIS BOOK BRIDGES THE GAP BETWEEN NEURO-CLINICIANS, NEURO-IMAGERS AND NEURO-PATHOLOGISTS, AND ALL ALLIED CLINICAL DISCIPLINES.
- DISEASES MARIE-FRANCOISE CHESSELET, 2000-10-19 WITH THE UNPRECEDENTED IDENTIFICATION OF NEW MUTATION MECHANISMS IN NEURODEGENERATIVE DISEASES AND THE EMERGENCE OF COMMON MECHANISMS AMONG DISEASES THAT WERE ONCE CONSIDERED UNRELATED, NEUROBIOLOGISTS ARE POISED FOR THE DEVELOPMENT OF NEW THERAPIES BASED ON HIGH THROUGHPUT SCREENINGS AND A BETTER UNDERSTANDING OF THE MOLECULAR AND CELLULAR MECHANISMS LEADING TO NEURODEGENERATION. IN MOLECULAR MECHANISMS OF NEURODEGENERATIVE DISEASES, MARIE-FRANCOISE CHESSELET, MD, PhD, AND A PANEL OF LEADING RESEARCHERS AND NEUROLOGISTS FROM INDUSTRY AND ACADEMIA CRITICALLY REVIEW THE MOST RECENT ADVANCES FROM DIFFERENT YET COMPLEMENTARY POINTS OF VIEW. FOCUSING ON ALZHEIMER'S, PARKINSON'S, AND CAG TRIPLET REPEAT DISEASES, THE AUTHORS SHOW HOW STUDIES OF CELLULAR AND GENETICALLY ENGINEERED ANIMAL MODELS HAVE ENHANCED OUR UNDERSTANDING OF THE MOLECULAR MECHANISMS OF NEURODEGENERATIVE DISEASES AND MAY LEAD TO THE DEVELOPMENT OF NEW THERAPEUTICS. TOPICS INCLUDE THE ROLE OF AB TOXICITY, GLIAL CELLS, AND INFLAMMATION IN ALZHEIMER'S DISEASE; THE FORMATION OF ABNORMAL PROTEIN FRAGMENTS ACROSS SEVERAL DISEASES, THE IMPACT OF DOPAMINE AND MITOCHONDRIAL DYSFUNCTION ON NEURODEGENERATION; AND THE POTENTIAL OF GENETICS TO IDENTIFY THE MOLECULAR MECHANISMS OF NEURODEGENERATIVE DISEASES. AUTHORITATIVE AND INSIGHTFUL, MOLECULAR MECHANISMS OF NEURODEGENERATIVE DISEASES SYNTHESIZES THE NOVEL IDEAS AND CONCEPTS NOW EMERGING TO CREATE A FRESH UNDERSTANDING OF NEURODEGENERATIVE DISOADERS, ONE THAT PROMISES TO LEAD TO POWERFUL NEW THERAPIES THAT PREVENT, DELAY THE ONSET, SLOW THE PROGRESSION, OR EVEN CURE THESE CRUEL DISEASES.
- DOCHEMISTRY AND CELL BIOLOGY OF AGEING: PART II CLINICAL SCIENCE J. ROBIN HARRIS, VIKTOR I. KOROLCHUK, 2019-03-19 THIS VOLUME OF THE SUBCELLULAR BIOCHEMISTRY SERIES WILL ATTEMPT TO BRIDGE THE GAP BETWEEN THE SUBCELLULAR EVENTS THAT ARE RELATED TO AGING AS THEY WERE DESCRIBED IN THE FIRST VOLUME OF THIS SET OF TWO BOOKS AND THE REALITY OF AGING AS THIS IS SEEN IN CLINICAL PRACTICE. ALL CHAPTERS WILL START FROM THE BIOCHEMISTRY OR CELL BIOLOGY, WHERE THE DATA IS AVAILABLE AND WORK UP TOWARDS THE UNDERSTANDING THAT WE HAVE OF AGING IN THE VARIOUS AREAS THAT ARE RELATED TO THE SUBJECT. KEY FOCUS POINTS FOR THIS VOLUME ARE NUTRITION, EXTERNAL FACTORS AND GENETICS ON AGING. THERE WILL ALSO BE CHAPTERS THAT WILL FOCUS ON VARIOUS ORGANS OR TISSUES IN WHICH AGING HAS BEEN WELL STUDIED, LIKE THE EYES, THE MUSCLES, THE IMMUNE SYSTEM AND THE BOOKS. THE AIM OF THE BOOK PROJECT AND THE BOOK PROJECT THAT IS PUBLISHED IN CONCERT WITH THIS VOLUME IS TO BRING THE SUBCELLULAR AND CLINICAL AREAS INTO CLOSER CONTACT.
- DECEMBERATIVE DISEASES PHILIP BEART, MICHAEL ROBINSON, MARCUS RATTRAY, NICHOLAS J. MARAGAKIS, 2017-07-03 PROVIDES A TIMELY OVERVIEW OF CRITICAL ADVANCES IN MOLECULAR AND CELLULAR NEUROBIOLOGY, COVERS KEY METHODOLOGIES DRIVING PROGRESS, AND HIGHLIGHTS KEY FUTURE DIRECTIONS FOR RESEARCH ON NEURONAL INJURY AND NEURODEGENERATION RELEVANT TO NEURONAL BRAIN PATHOLOGIES. THE EDITORS BRING TOGETHER CONTRIBUTIONS FROM INTERNATIONALLY RECOGNIZED WORKERS IN THE FIELD TO PROVIDE AN UP TO DATE ACCOUNT OF HOW AND WHY MOLECULAR AND CELLULAR NEUROBIOLOGY IS SUCH AN IMPORTANT AREA FOR CLINICAL NEUROSCIENCE. UNDERSTANDING THE MOLECULAR ASPECTS OF A NUMBER OF NEURODEGENERATIVE CONDITIONS SUCH AS PARKINSON'S OR ALZHEIMER'S DISEASE FOR THE PURPOSE OF IMPROVING PATIENT MANAGEMENT REMAINS A MAJOR CHALLENGE OF NEUROBIOLOGY BE IT FROM THE BASIC OR CLINICAL PERSPECTIVE. A STRATEGIC EVALUATION OF RESEARCH CONTRIBUTIONS AND THE POWER OF MODERN METHODS WILL HELP ADVANCE KNOWLEDGE OVER THE NEXT YEARS.

- DEMENTIA WITH LEWY BODIES JOHN O'BRIEN, DAVID AMES, IAN MCKEITH, EDMOND CHIU, 2005-11-29 FILLING A NOTICEABLE GAP IN THE MARKET FOR A NEW TEXT SOLELY FOCUSED ON DEMENTIA WITH LEWY BODIES, THIS BOOK DISCUSSES CUTTING-EDGE TOPICS COVERING THE CONDITION FROM DIAGNOSIS TO MANAGEMENT, AS WELL AS WHAT IS KNOWN ABOUT THE NEUROBIOLOGICAL CHANGES INVOLVED. WITH HUGE PROGRESS HAVING BEEN MADE OVER THE LAST DECADE IN TERMS OF THE DISORDER
- Pathology, Prevention and Therapeutics of Neurodegenerative Disease Sarika Singh, Neeraj Joshi, 2018-09-24 This book, written by a leading panel of experts in the field of neurosciences, provides a comprehensive overview of the pathology of neurodegenerative diseases as well as the preventive measures. Prevention is important due to the lack of early diagnostic markers and the limitations/ problems of treating neurodegenerative diseases
- MP-activated Protein Kinase Mario D. Cordero, Benoit Viollet, 2016-11-02 AMPK has emerged as an important integrator of signals that control energy balance through the regulation of multiple biochemical pathways in eukaryotes. This book focuses on the implications of AMPK as a master metabolic regulator in diseases, including new methods and animal models. The contributions are written by leading experts in the field and give an extensive overview of the current knowledge of AMPK biology and the role of AMPK in health and disease.
- Eminciples and Practice of Movement Disorders E-Book Joseph Jankovic, Mark Hallett, Stanley Fahn, 2011-04-01 Principles and Practice of Movement Disorders provides the complete, expert guidance you need to diagnose and manage these challenging conditions. Drs. Stanley Fahn, Joseph Jankovic and Mark Hallett explore all facets of these disorders, including the latest rating scales for clinical research, neurochemistry, clinical pharmacology, genetics, clinical trials, and experimental therapeutics. This edition features many new full-color images, additional coverage of pediatric disorders, updated Parkinson information, and many other valuable updates. An accompanying Expert Consult website makes the content fully searchable and contains several hundred video clips that illustrate the manifestations of all the movement disorders in the book along with their differential diagnoses. Get just the information you need for a clinical approach to diagnosis and management, with minimal emphasis on basic science. Find the answers you need quickly and easily thanks to a reader-friendly full-color format, with plentiful diagrams, photographs, and tables. Apply the latest advances to diagnosis and treatment of pediatric movement disorders, Parkinson disease, and much more. View the characteristic presentation of each disorder with a complete collection of professional-quality, narrated videos online. Better visualize every concept with new full-color illustrations throughout. Search the complete text online, follow links to PubMed abstracts, and download all of the illustrations, at www.expertconsult.com.
- EMEW PROGNOSTIC AND PREDICTIVE MARKERS IN CANCER PROGRESSION SUSAN COSTANTINI ALFREDO BUDILLON, ALFREDO BUDILLON, 2021-02-12 BIOMARKERS ARE OF CRITICAL MEDICAL IMPORTANCE FOR ONCOLOGISTS, ALLOWING THEM TO PREDICT AND DETECT DISEASE AND TO DETERMINE THE BEST COURSE OF ACTION FOR CANCER PATIENT CARE. PROGNOSTIC MARKERS ARE USED TO EVALUATE A PATIENT'S OUTCOME AND CANCER RECURRENCE PROBABILITY AFTER INITIAL INTERVENTIONS SUCH AS SURGERY OR DRUG TREATMENTS AND, HENCE, TO SELECT FOLLOW-UP AND FURTHER TREATMENT STRATEGIES. ON THE OTHER HAND, PREDICTIVE MARKERS ARE INCREASINGLY BEING USED TO EVALUATE THE PROBABILITY OF BENEFIT FROM CLINICAL INTERVENTION(S), DRIVING PERSONALIZED MEDICINE. EVOLVING TECHNOLOGIES AND THE INCREASING AVAILABILITY OF "MULTIOMICS" DATA ARE LEADING TO THE SELECTION OF NUMEROUS POTENTIAL BIOMARKERS, BASED ON DNA, RNA, MIRNA, PROTEIN, AND METABOLIC ALTERATIONS WITHIN CANCER CELLS OR TUMOR MICROENVIRONMENT, THAT MAY BE COMBINED WITH CLINICAL AND PATHOLOGICAL DATA TO GREATLY IMPROVE THE PREDICTION OF BOTH CANCER PROGRESSION AND THERAPEUTIC TREATMENT RESPONSES. HOWEVER, IN RECENT YEARS, FEW BIOMARKERS HAVE PROGRESSED FROM DISCOVERY TO BECOME VALIDATED TOOLS TO BE USED IN CLINICAL PRACTICE. THIS SPECIAL ISSUE COMPRISES EIGHT REVIEW ARTICLES AND FIVE ORIGINAL STUDIES ON NOVEL POTENTIAL PROGNOSTIC AND PREDICTIVE MARKERS FOR DIFFERENT CANCER TYPES.
- D-1/PARK7 PROTEIN HIROYOSHI ARIGA, SANAE M. M. IGUCHI-ARIGA, 2017-11-16 THIS BOOK REVIEWS THE FUNCTIONS AND ROLES OF DJ-1 IN VARIOUS OXIDATIVE STRESS-RELATED DISEASES AND APPLICATIONS OF DJ-1 AND ITS BINDING COMPOUNDS TO THE DISEASES. THE DJ-1 GENE WAS FIRST FOUND TO BE A NOVEL ONCOGENE IN 1997 AND LATER, IN 2003, ALSO FOUND TO BE A CAUSATIVE GENE FOR A FAMILIAL FORM OF PARKINSON'S DISEASE (PD), PARK7. THE DJ-1 GENE IS THEREFORE THE FIRST GENE DISCOVERED THAT IS KNOWN TO CAUSE CANCER AND NEURODENEGERATIVE DISEASES, INCLUDING PD. THE RESEARCH FIELD HAS EXPANDED AS THE RESEARCH HAS DEVELOPED. THUS THIS VOLUME BEGINS WITH A GENERAL INTRODUCTION OF DJ-1, AND EXPLAINS THE HISTORY AND RESEARCH DEVELOPMENT TO UNDERSTAND THE FOLLOWING CHAPTERS. THOSE CHAPTERS PRESENT THE ROLES OF DJ-1 IN VARIOUS OXIDATIVE STRESS-RELATED DISEASES SUCH AS NEURODEGENERATIVE DISEASES, AS WELL AS CANCER, DIABETES, AND FERTILITY. MOREOVER, SEVERAL CHAPTERS PRESENT EVIDENCE THAT DJ-1 IS USEFUL FOR THERAPEUTIC STRATEGIES AGAINST THESE DISEASES. THE READER WILL DISCOVER THAT DJ-1 IS A PROMISING PROTEIN BOTH FOR BASIC CELL BIOLOGY AND FOR THE MECHANISM AND THERAPY FOR OXIDATIVE STRESS-RELATED DISEASES.

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