



Target World Series Review

L Cohen



Target World Series Review:

Novel Therapeutic Target and Drug Discovery for Neurological Diseases, Volume II Rui Liu, Zhuorong Li, George Barreto, Yong Cheng, Liu Qing-Shan, 2025-03-12 This Research Topic is the second volume of the Research Topic Novel Therapeutic Target and Drug Discovery for Neurological Diseases Please see the first volume here Neurological diseases represent heterogeneous disorders that involve complex pathological alterations with only a few effective treatments therefore there is a great need for the development of novel therapeutic targets and therapies There are more than six hundred neurological disorders including dementia cerebrovascular diseases mental illness brain tumor and infection caused central diseases all arising from disruption to the central nervous system Researchers are attempting to identify promising targets and potential drug candidates from new biological techniques and pharmacological animal studies Thus novel target discovery and mechanism elucidation as well as the development of small molecule entities natural compounds and polypeptides for the treatment of neurological dysfunction may offer exciting clues for novel therapeutic strategies to halt or mitigate the course of neurological diseases

Vaccines and Therapeutics Utilizing New Adjuvants and Potential Inhibitors to Target Emerging Infectious Diseases Jiae Kim, Ousman Jobe, Mara Jana Broadhurst, 2025-12-01 Emerging infectious diseases such as persistent infectious diseases that have not been controlled i.e. HIV dengue fever those increasing in incidence i.e. flaviviruses food and waterborne pathogens or previously unknown diseases have impacted populations throughout the world These diseases require either preventative vaccines or proper therapeutics to mitigate global epidemics from emerging Recent vaccine and therapeutic candidates have emerged and are moderately successful however there is still room for improvement with either long lasting vaccine efficacy or therapeutics with maximal efficacy and minimal toxicity New technologies are necessary for both preventative vaccines and therapeutics for emerging infectious diseases As these diseases impact populations on a global scale preventing infection is key However when that is not feasible treatment strategies and specific therapeutics with minimal toxicity and side effects are just as critical in fighting such epidemics This Research Topic aims to cover current and emerging technologies being explored to prevent or treat infectious diseases including strategies currently in preclinical or clinical phase studies We aim to bring together all aspects of research and development of these new technologies and invite vaccinologists immunologists pharmacologists and other multidisciplinary groups to feed knowledge into these growing fields This research will impact the current landscape of infectious diseases and may also provide interventions against future pandemics We welcome the submission of Review and Original Research articles focusing on preclinical and clinical vaccines or therapeutic studies and Perspective and Opinion articles highlighting the gaps in vaccines and therapeutics targeting emerging infectious diseases including viral or bacterial infections as well as multidrug resistant bacteria Articles may focus on but are not limited to the following subtopics Promising adjuvants in preclinical and clinical vaccine trials Promising inhibitors that target infectious diseases in preclinical studies Mechanism of

action of novel inhibitors or adjuvants Formulations that target different sites of action i e mucosal regions **Novel**

Approaches to Target the Immune System in Gastrointestinal Cancers Gianluigi Giannelli, Erica Villa, Maria L.

Martinez Chantar, 2022-01-31 *Target Organ Damage in Fabry Disease* Guido Iaccarino, Francesca Graziani, 2025-08-27

Anderson Fabry disease is a rare genetic disease characterized by a defect in a X Linked gene the alpha galactosidase Due to the sexual chromosome localization of the gene men patients present a more severe phenotype characterized by an early occurrence of kidney failure cardiac hypertrophy neurological disorders including cryptogenic stroke as well as pain hypo hidrosis skin lesions corneal depositions fatigue Women affected by this disease present a late onset phenotype that is characterized by a lighter symptomatology including fatigue and organ damage Also the number of mutations in the alpha galactosidase gene that could affect the enzyme activity or expression has raised up to over 1000 The association between genetic alterations and the phenotype severity is very much difficult to be identified *The Ischemic Penumbra: Still the*

Target for Stroke Therapies? Argye E. Hillis, Jean-Claude Baron, 2015-07-23 The ischemic penumbra was initially defined by Symon Lassen and colleagues in the 1970s as an area of brain tissue with inadequate blood flow to maintain electric activity of neurons but adequate blood flow to preserve the function of the ion channels This area of tissue receiving enough blood to survive but not enough to function often surrounds or abuts the irreversibly damaged core in ischemic stroke It was shown that if blood flow could be restored to this area of marginal perfusion the tissue could survive and function again and growth of the core could be prevented Based on seminal PET studies penumbra or penumbral tissue eventually took on a subtly different meaning the area of brain that is destined to progress to infarct unless blood flow is restored within a particular time window The penumbra thus became the target for all acute stroke interventions to preserve viability of the tissue and restore function New imaging techniques including diffusion and perfusion MRI and CT perfusion were developed to rapidly identify individuals with penumbra who were thought to be the best candidates for aggressive interventions to restore blood flow particularly beyond the licensed time window for IV thrombolysis However most clinical trials have failed to establish the usefulness of identifying candidates for treatment in this way using pre specified protocols and primary endpoints These trials have used different and sometimes unvalidated thresholds of hypoperfusion as well as irreversible infarct and various definitions of significant penumbra or mismatch between irreversible infarct and hypoperfused but salvageable tissue and reanalysis of their data using more refined image processing showed post hoc positivity They have also evaluated outcome in a variety of ways with few studies measuring the direct effect of restoring blood flow on the function of the penumbral tissue Therefore important remaining questions include how to define characterize and image the penumbra in acute stroke to achieve the greatest reliability and validity for what we want to measure and whether this concept so defined provides an optimal target for stroke therapy using state of the art trial design *Precision Immunotherapy and Novel Target Discovery*

in Hematological Malignancy Beibei Zhang, Tuo Liu, Yu Pan, 2025-11-13 Hematological malignancies represent defying

clinical conditions affecting either the lymphoid or the myeloid lineages Targeted therapy research is developing rapidly from bench to bedside based on the continually up dated disease molecular understanding and growing novel target identifications in malignancy Patient survival expectations and clinical outcomes are closely associated with the development of precision treatments For example acute myeloid leukemia AML is a group of myeloid malignancies characterized by high heterogeneity in clinical courses and therapy responses The emergence of new targeted agents such as FLT3 BCL2 inhibitors or immune checkpoint blockade ICB therapy has significantly improved the survival of a subtype of AML In addition inspired by the promising outcomes of CAR T therapy in certain malignancies various new immunotherapy technologies are emerging rapidly based on neoantigens Nevertheless the lack of ideal targets or neoantigens seems to be one of the major stumbling blocks in the precision immunotherapy of most hematological malignancies Currently developed technological innovations have driven the identification of more and more malignant disease specific novel molecules or neoantigens that arise as diagnostic biomarkers and therapeutic targets Thus precision targeted immunotherapy is developing as one of the most promising approaches in the battle against these hematological malignant diseases There is no doubt that the development of novel precision diagnoses and therapeutic approaches against these novel targets or neoantigens will outline the prospective clinical applications in the future Given the significant clinical outcomes of novel target based immunotherapy and uncovered mechanism understanding in hematological malignancies the research topic welcomes studies involved in the latest developments in novel target or neoantigen discoveries and utilizations including small chemical molecular immunomodulatory drug development vaccines antibodies and cell therapies from basic preclinical research to translational studies in hematological malignancies The main goal of this research topic is to draw a clear framework within which we can see the steps that should be taken to establish and maintain effective anti malignancy immune responses based on novel target or neoantigen associated immunotherapeutic approaches which are crucial for interpreting hematological malignancy clinical outcomes The research topic welcomes submissions of original research articles brief research reports clinical cases perspectives mini reviews and reviews concerning hematological malignancies Topics include but are not limited to 1 Novel target or neoantigen screening and discovery method development 2 Novel target or neoantigen based malignancy diagnosis 3 Novel small chemical molecular immunomodulatory drug development 4 Novel vaccine antibody development and cell immunotherapy 5 Immune escape or resistance mechanisms Please note manuscripts consisting solely of bioinformatics or computational analysis of public genomic or transcriptomic databases which are not accompanied by validation independent cohort or biological validation in vitro or in vivo are out of scope for this section and will not be accepted as part of this Research Topic

Immune Cell Interactions With Target Cells in Physiological and Pathological Conditions of the Nervous System Antje Kroner, Sven G. Meuth, Craig Stephen Moore, 2020-09-02 This eBook is a collection of articles from a Frontiers Research Topic Frontiers Research Topics are very popular trademarks of the Frontiers Journals Series they are collections

of at least ten articles all centered on a particular subject With their unique mix of varied contributions from Original Research to Review Articles Frontiers Research Topics unify the most influential researchers the latest key findings and historical advances in a hot research area Find out more on how to host your own Frontiers Research Topic or contribute to one as an author by contacting the Frontiers Editorial Office frontiersin.org/about/contact Computational and Experimental Approaches in Multi-target Pharmacology Thomas J. Anastasio, 2017-08-24 The next frontier in pharmacology is the development of multi target strategies in which pathological processes are controlled by pharmacologically manipulating them at many different points at once Designing multi target strategies will require deep understanding of the complex physiology that underlies pathological processes It will also require the development of single drugs with multiple targets or combinations of drugs with compatible pharmacokinetics that work synergistically to maximize desirable effects while minimizing unwanted side effects This e Book contains ten original articles each addressing a different aspect of this challenge Together they open new perspectives and show the way forward in the development of multi target therapeutics

Inhibiting PARP as a Strategic Target in Cancer Christina Annunziata, Kristin Zorn, 2016-08-05 Poly ADP ribose polymerase PARP proteins are critical mediators of DNA repair Many traditional anti cancer chemotherapy agents overwhelm a cell's ability to repair DNA damage in order to kill proliferating malignant cells Recent evidence suggests that cancers within and across tissue types have specific defects in DNA repair pathways and that these defects may predispose for sensitivity and resistance to various classes of cytotoxic agents Breast ovarian and other cancers develop in the setting of inherited DNA repair deficiency and these cancers may be more sensitive to cytotoxic agents that induce DNA strand breaks as well as to inhibitors of PARP activity A series of recent clinical trials has tested whether PARP inhibitors can achieve synthetic lethality in hereditary DNA repair deficient tumors At the current time mutation of BRCA serves as a potential but not comprehensive biomarker to predict response to PARP inhibitor therapy Mechanisms of resistance to PARP inhibitors are only recently being uncovered Future studies seek to identify sporadic cancers that harbor genomic instability rendering susceptibility to PARP inhibitors that compound lethal DNA damage

Evolution of NK-mediated target recognition under the pressure of physiologic or pathologic stimuli Massimo Vitale, Simona Sivori, Miguel López-Botet, Daniel Olive, 2015-03-23 Since their discovery NK cells have come out as potential tools to fight cancer and viruses This finding early urged different groups to study the mechanisms governing NK cell function The identification of the MHC I specific inhibitory receptors i.e. KIRs NKG2A and certain Ly49 molecules allowed defining rather rapidly how NK cells could avoid self aggression and how they could be directed towards targets that were forced by viral infection or tumor transformation to down regulate MHC I expression In a second time also the repertoire of surface activating receptors addressing NK cytotoxicity towards tumors and pathogens was mostly defined In spite of the first findings however most recent studies may suggest that NK cells and their receptors might not have been evolved to kill tumor targets and perhaps they might have

been only partially influenced in their evolution by the need of recognizing viruses Indeed certain NK receptors known to activate NK cell cytotoxicity NKp30 DNAM 1 NKp80 can also participate at regulatory interactions occurring between NK and myeloid cells In addition a peculiar NK cell subset which intensively populate decidua during the first trimester of pregnancy through the engagement of specific receptors and the interaction with decidual DC produce chemokines and pro angiogenic cytokines and induce Tregs Thus in this context NK cells favor decidua vascularization and development of the semiallogeneic foetus in a tolerant environment Viruses have nevertheless played an important role in shaping the NK cell receptor repertoire Several studies have unveiled clues of the evolutionary struggle between these pathogens and NK cells Different NK receptors including NKp46 NKp30 NKp44 NKG2D NKG2C Ly49 and certain KIRs have been demonstrated to recognize virus encoded or virus induced ligands The expression of TLR specifically recognizing microbial products together with the unexpected role of KIR3DL2 in shuttling these products to TLR containing endosomes have also been documented in NK cells On the other side different viral immune evasion molecules have been shown to interfere with the expression of ligands for T or NK cell activating receptors In addition viral infections can occur in the reproductive stage of life cycle and may represent a serious threat for the species propagation Thus the control of viruses together with the maintenance of foetus during pregnancy should represent major evolutionary forces in shaping NK receptors Along this line the NK mediated control of tumors should not be under the same evolutionary pressure as tumors mostly appear later in the life cycle and the recognition of tumor encoded ligands may be less efficient as the NK cell receptors might have not been selected for such aim This may be the reason why although displaying strong antitumor activity in vitro NK cells could hardly contain tumor burden in vivo In addition the pathogen driven evolution of NK cell function may also favor the role of NK cells in the insurgence of immune mediated diseases This research topic will collect contributions that may clarify the relationships between the evolution of the NK receptors and their role in an efficient recognition of viruses and tumor cells or in immune mediated diseases

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