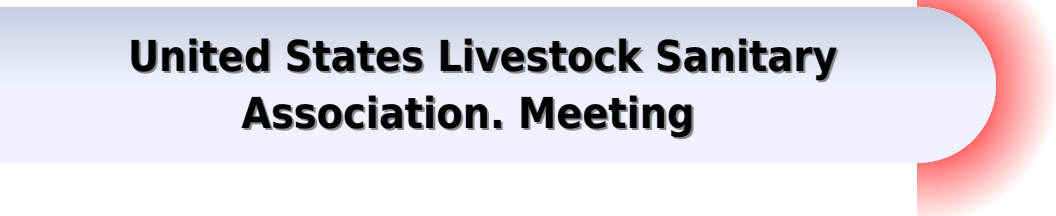




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Innovative Therapeutic and Vaccine Approaches against Respiratory Pathogens Christophe Chevalier, Cynthia Calzas, Delphyne Descamps, Michel Chignard, 2020-05-06 Challenges and Opportunities for Respiratory Syncytial Virus Vaccines Larry J. Anderson, Barney S. Graham, 2013-12-20 Although respiratory syncytial virus RSV has been a high priority for vaccine development for over 50 years now still no vaccine is available and none has yet demonstrated sufficient promise to move to licensure The success of RSV immune prophylaxis and the availability of ever more powerful tools to study the immune response and pathogenesis of disease combined with the ability to construct a wide variety of vaccines using different vaccine platforms give us grounds to believe that an RSV vaccine is within reach This book brings together in one source what is currently known about the virus its clinical and epidemiologic features the host response and pathogenesis of the disease vaccines vaccine platforms and treatment and animal and tissue culture models of RSV infection It is designed to organize the critical information relevant to RSV vaccine development facilitate the assimilation of data and speed progress toward producing a safe and effective vaccine *Current Progress and Challenges in the Development of a Hepatitis C Virus Vaccine* Steven K.H. Fong, Thomas F. Baumert, 2019-01-22 More than 70 million people worldwide are infected with hepatitis C virus a major cause of liver cirrhosis liver failure and hepatocellular carcinoma world wide In the last decade this cancer has emerged as the second leading cause of cancer death and the global burden is increasing by two million new infections per year mainly due to injection drug use An effective vaccine will be the most effective means to contain the spread of this virus worldwide The articles in this Research Topic describe the progress that has been made towards a preventive vaccine and the challenges that still need to be overcome to ultimately achieve this goal The United States-Japan Cooperative Medical Science Program, 1990 Viral Infection at the Maternal-Fetal Interface Abhay P. S. Rathore, Vivian Vasconcelos Costa, Ashley L. St John, 2022-02-28 **Dengue Virus-Specific T Cell Immunity** Laura Rivino, Gathsaurie Neelika Malavige, Daniela Weiskopf, 2020-08-05 *Influenza Virus Vaccines and Immunotherapies* Arun Kumar, Shakti Singh, 2016-03-15 Influenza virus infections lead to thousands of deaths worldwide annually and billions of dollars economic burden Despite continuing advances in our understanding of the immune evasion mechanism the disease remains one of the foremost threat for human being Traditional vaccines attenuated and inactivated mainly provide protection by inducing virus neutralizing antibodies targeting ever changing surface antigens Haemagglutinin HA and Neuraminidase NA Due to genetic shift and immune selection pressure prevalence of circulating influenza virus subtypes changes every year Therefore mismatch between circulating strain and vaccine strain can critically affect the success rate of these conventional flu vaccines and requires continuous monitoring of circulating influenza virus subtypes and change in the vaccine formulations accordingly The collective limitations of existing flu vaccines urgently call for the development of a novel universal vaccines that might provide the required protective immunity to a range of influenza virus subtypes New

approaches are being investigated mainly targeting conserved regions of flu proteins. Some of these approaches include universally conserved epitopes of HA nucleoprotein NP, capsid protein M1, and ion channel protein M2 that induced strong immune responses in animal models. Some attention and progress appears to be focused on vaccines based on the M2 ectodomain M2e employing a variety of constructs, adjuvants, and delivery systems including M2e hepatitis B core antigen, flagellin constructs, and virus like particles (VLP). Animal studies with these M2e candidate vaccines demonstrated that these vaccine candidates can prevent severe illness and death but not infection, which may pose difficulties in both the evaluation of clinical efficacy and approval by the regulatory authorities. VLP vaccines appear to be promising but still are mostly limited to animal studies. The discovery and development of new and improved vaccines have been greatly facilitated by the application of new technologies. The use of nucleic acid based vaccines to combine the benefits of in situ expression of antigens with the safety of inactivated and subunit vaccines has been a key advancement. Upon their discovery more than 20 years ago, nucleic acid vaccines promised to be a safe and effective means to mimic immunization with a live organism vaccine, particularly for induction of T cell immunity. In addition, the manufacturing of nucleic acid based vaccines offered the potential to be relatively simple, inexpensive, and generic. Reverse vaccinology and in silico designing of vaccines are very innovative approaches and are being considered as the future of vaccines. Furthermore, various immuno therapeutic agents also are being developed to treat and minimize immuno pathological damage in patients suffering from life threatening complications. For the treatment of such pathological conditions, various novel approaches such as administration of immune suppressive cytokines, blocking co stimulatory signals, or activating co inhibitory signals of T cell activation are being tested both in lab and clinics. The Research Topic on influenza virus vaccine and therapeutics will give an insight into the current status and future scope of these new innovative approaches and technologies. Moreover, these new methods will also serve as a reference tool for the development of future vaccines against several other pathogens.

Why Vaccines to HIV, HCV and Malaria Have So Far Failed - Challenges to Developing Vaccines against Immunoregulating Pathogens Shuo Li, Magdalena

Plebanski, Peter Smooker, Eric J. Gowans, 2016-09-14. Despite continuous progress in the development of anti viral and anti bacterial parasite drugs, the high cost of medicines and the potential for re infection, especially in high risk groups, suggest that protective vaccines to some of the most dangerous persistent infections are still highly desirable. There are no vaccines available for HIV, HCV, and Malaria, and all attempts to make a broadly effective vaccine have failed so far. In this Research Topic, we look into why vaccines have failed over the years and what we have learned from these attempts. Rather than only showing positive results, this issue aims to reflect on failed efforts in vaccine development. Coming to understand our limitations will have theoretical and practical implications for the future development of vaccines to these major global disease burdens.

Proceedings of the National Academy of Sciences of the United States of America National Academy of Sciences (U.S.), 2007 *Proceedings, Annual Meeting of the United States Livestock Sanitary Association* United States

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