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Jesse Goodman, M.D., M.P.H. is Professor of Medicine at Georgetown University and Director of the Center on Medical Product Access, Safety and Stewardship, focusing on science and policy to address public health needs. He is an active clinician and Attending Physician in Infectious Diseases at Georgetown University, Washington DC Veterans Administration and Walter Reed Medical Centers. Until 2014, he was Chief Scientist of the U.S. Food and Drug Administration (FDA), also serving as part of the government's senior leadership for the 2009 influenza pandemic and other major public health responses as well as the 2010 White House Medical Countermeasure Review. Prior to that, he directed FDA's Center for Biologics Evaluation and Research (CBER) and before that, as Senior Advisor to the FDA Commissioner, developed and co-chaired the U.S. Task Force to Combat AMR, producing the first National Action Plan. Previously he was Professor of Medicine and Chief of Infectious Diseases at the University of Minnesota, where his laboratory isolated *A. phagocytophilum*, the causative agent of human granulocytic anaplasmosis. He has served on various CDC, NIH, DOD, CEPI, National Academy of Medicine (NAM) and WHO Advisory Committees, including the WHO SAGE for Immunization and Ebola Vaccine Working Group and, recently, the NAM Consensus Committee on Lyme Infection Associated Chronic Illness. He co-Chairs the multi-institution COVID Vaccine Analysis Team (COVAT), now E-VAT (Expert Vaccine Analysis Team). He is on the Boards of the US Pharmacopeia, GSK (through May, 2025), Intellia Therapeutics (chairing its Science and Technology committee) and BiomX. He has been elected to the National Academy of Medicine of the National Academy of Sciences.