

## NICHOLAS A. MEANWELL, Ph.D., CSci, CChem, FRSC

### EDUCATION

**B.Sc. Special Honours Chemistry, 1976**  
The University of Sheffield, Sheffield, England

**Ph.D. Degree, Organic Chemistry, 1979**  
The University of Sheffield, Sheffield, England  
Under the supervision of Dr. D. Neville Jones

**Post Doctoral Studies, 1979-1982**  
Wayne State University, Detroit, Michigan  
In collaboration with Professor Carl R. Johnson

### INDUSTRIAL EXPERIENCE

Bristol-Myers Squibb Research and Development: 1982-present

**Current Position:** Executive Director, Department of Discovery Chemistry, Virology

Have led drug discovery programs in the cardiovascular, neurosciences and virology therapeutic areas, work that has resulted in the advancement of over 25 clinical candidates for the prevention of thrombosis, the treatment of stroke and therapy for viral infections, including human immunodeficiency virus (HIV), hepatitis C virus (HCV) and respiratory syncytial virus (RSV). Recent significant discoveries to emerge from this group include RSV fusion inhibitors, characterized as the first small molecules to interfere with the association of the 6 helical peptide bundle that is a critical step in the virus entry process, and a series of HIV attachment inhibitors that are the first small molecules described to function by interfering with the interaction between virus gp120 and the host cell CD4 receptor. BMS-663068, a phosphonooxymethyl prodrug of BMS-626529, that has just successfully completed Phase 2 clinical trials.

Significant compounds in the HCV arena include daclatasvir, a pioneering molecule that established NS5A inhibition as a clinically-relevant target and which has completed Phase 3 clinical trials, the HCV NS3 protease inhibitor asunaprevir which incorporates the cyclopropyl acylsulfonamide moiety that has been widely adopted, and BMS-791325, a thumb site inhibitor of HCV NS5B polymerase which is currently in Phase 3 trials. The combination of daclatasvir and asunaprevir in a clinical trial conducted in HCV-infected subjects established for the first time that a HCV infection could be cured by direct acting antiviral agents in the absence of immune stimulation. NDAs seeking marketing approval for the combination of daclatasvir and asunaprevir have been filed in Japan and the United States.

### PUBLICATIONS and PATENTS

Author/co-author of 140 peer-reviewed publications, 45 review articles and book chapters and more than 150 meeting abstracts. Named as inventor/co-inventor of over 100 issued U.S. Patents and have presented over 65 invited lectures at National and International meetings, Universities and Schools on Medicinal Chemistry.

