CURRICULUM VITAE

Last Name:	Cook	**First Name:**	Jack	**Middle Initial:**	A	**Today's Date:**	18 June 2018

Job Title:	Vice President, Clinical Pharmacology TA Head Rare Disease & INI
Professional Address:	Global Innovative Pharma Business Unit, Pfizer Inc 445 Eastern Point Road Groton, CT 06340

Educational History (Relevant Academic and/or Professional Qualifications)

Degree/Certification, Field	Date	Institution, Country
Ph.D., Pharmaceutics	Sep 1987	University of Michigan, USA
B.S., Pharmacy	May 1981	Ferris State College, USA
B.S., Applied Mathematics	May 1981	Ferris State College, USA
A.A.S. Industrial Chemical Technology	May 1978	Ferris State College, USA

Current Position and Previous Relevant Positions Held (most current data first)

2016–present, Vice President, GPD, Clinical Pharmacology, Pfizer Inc., USA

- Provides line leadership and management to Clinical Pharmacology Leads.
- Responsible for resourcing, recruitment, supervision, training and staff development in clinical pharmacology. Coaches and develops Clinical Pharmacology Leads.
- Accountable and responsible for the clinical pharmacology deliverables for the assigned projects within sphere of influence either directly or via CPLs.
- Responsible for ensuring appropriate Clinical Pharmacology representation on drug development and clinical teams and provision of clinical pharmacology expertise and leadership to a project.
- Responsible for ensuring planning and direction of clinical pharmacology components of clinical programs (including clinical development plan/life cycle plan) and studies (including synopsis preparation; clinical phase oversight, reporting).
- Responsible for implementing clinical pharmacology best practices either directly or via Clinical Pharmacology Leads.

2014-2016, Vice President, Global Innovative Pharma, Clinical Pharmacology, Pfizer Inc., USA

- Provides line leadership and management to Clinical Pharmacology Leads.
- Responsible for resourcing, recruitment, supervision, training and staff development in clinical pharmacology. Coaches and develops Clinical Pharmacology Leads.
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- Responsible for implementing clinical pharmacology best practices either directly or via Clinical Pharmacology Leads.

2009-2013, Vice President, Specialty Care, Clinical Pharmacology, Pfizer Inc., USA

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- Responsible for implementing clinical pharmacology best practices either directly or via Clinical Pharmacology Leads.

2008-2009, Executive Director, Global R&D, Clinical Pharmacology, Pfizer Inc., USA
- Provides line leadership and management to Clinical Pharmacology Leads.
- Responsible for resourcing, recruitment, supervision, training and staff development in clinical pharmacology. Coaches and develops Clinical Pharmacology Leads.
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- Responsible for implementing clinical pharmacology best practices either directly or via Clinical Pharmacology Leads.

2010-Present, Adjunct Faculty, College of Pharmacy, University of Florida, USA
- Lecturer in the area of drug development

2009-Present, Adjunct Professor of Pharmaceutical Sciences, University of Michigan, USA
- Lecturer in the area of drug development

2004-2009, Adjunct Associate Professor of Pharmaceutical Sciences, University of Michigan, USA
- Lecturer in the area of drug development

1989-1990, Adjunct Instructor, Albany College of Pharmacy, USA
- Lecturer in the area of drug development

2004-2008, Senior Director, Global R&D, Clinical Pharmacology, Pfizer Inc., USA
- Responsible for ensuring planning and direction of clinical pharmacology components of clinical programs (including clinical development plan/life cycle plan) and studies (including synopsis preparation; clinical phase oversight, reporting).
- Responsible for implementing clinical pharmacology best practices

2001-2004, Director, Global R&D, Clinical Pharmacology, Pfizer Inc., USA
- Responsible for ensuring planning and direction of clinical pharmacology components of clinical programs (including clinical development plan/life cycle plan) and studies (including synopsis preparation; clinical phase oversight, reporting).
- Responsible for implementing clinical pharmacology best practices

1994-2001, Senior Research Associate, PDM, Parke-Davis, USA
- Responsible for ensuring planning and direction of clinical pharmacology components of clinical programs (including clinical development plan/life cycle plan) and studies (including synopsis preparation; clinical phase oversight, reporting).
- Responsible for implementing clinical pharmacology best practices

1992-1994, Research Associate, Pharmacokinetics, PDM, Parke-Davis, USA
- Responsible for ensuring planning and direction of clinical pharmacology components of clinical programs (including clinical development plan/life cycle plan) and studies (including synopsis preparation; clinical phase oversight, reporting).
- Responsible for implementing clinical pharmacology best practices

1990-1992, Senior Scientist, PDM, Parke-Davis, USA
- Responsible for ensuring planning and direction of clinical pharmacology components of clinical

programs (including clinical development plan/life cycle plan) and studies (including synopsis preparation; clinical phase oversight, reporting).
- Responsible for implementing clinical pharmacology best practices

1989-1990, Group Leader, Drug Metabolism, Sterling Drug Inc., USA
- Responsible for ensuring planning and direction of clinical pharmacology components of clinical programs (including clinical development plan/life cycle plan) and studies (including synopsis preparation; clinical phase oversight, reporting).
- Responsible for implementing clinical pharmacology best practices

1987-1989, Senior Research Scientist, Drug Metabolism, Sterling Drug Inc., USA
- Responsible for ensuring planning and direction of clinical pharmacology components of clinical programs (including clinical development plan/life cycle plan) and studies (including synopsis preparation; clinical phase oversight, reporting).
- Responsible for implementing clinical pharmacology best practices