## ADRIAN POWELL THOMAS, MD, FRACP, MRACMA

### INDUSTRY EXPERIENCE

Janssen, the Pharmaceutical Companies of Johnson & Johnson Raritan, NJ, USA

Head, Global Public Health2013-CurrentVice President and Global Head, Global Commercial Strategy Organization2011-CurrentPresident and Company Officer, Janssen Global Services LLC2011-Current

 Global responsibility for the cluster of commercial and scientific functions supporting Janssen, including Global Medical Affairs; Global Health Economics; Global Pharmaceutical Pricing; Global Strategic Analytics; Strategic Marketing Excellence; Global Access Policy; and Patient Reported Outcomes.

### World Wide Vice President, Market Access

2009-2011

Global responsibility for Health Economics, Pharmaceutical Pricing, and Patient Reported Outcomes.

### Global Head, Benefit Risk Management & Chief Safety Officer, Pharmaceuticals

2007-2009

• Global responsibility for the benefit-risk management of J&J pharmaceutical products.

### Vice President, Safety Sciences, Informatics & Operations

2005-2007

 Responsibility for epidemiology, surveillance, safety systems and operational processing for J&J pharmaceutical products, both R&D and marketed.

### Head of Global Clinical Safety & Epidemiology, J&J Pharmaceutical R&D

2002-2005

Global responsibility for post-marketing surveillance and epidemiology for pharmaceutical products.

### Asia-Pacific Regional Head of Drug Safety & Surveillance, Janssen Research Foundation 2001-2002

Responsibility for Asia-Pacific Region & global safety for select biologic therapies.

### Schering-Plough Corporation

Kenilworth, NJ, USA

Australasian Medical Affairs Director, Schering-Plough

1999-2001

Responsibility for Medical Affairs functions in Australia and New Zealand.

### Eli Lilly & Company

West Ryde, New South Wales, Australia

Associate Medical Director, Australia & New Zealand	1998-1999
Associate Medical Director, Clinical Research	1997-1998
Clinical Research Physician, Eli Lilly	1995-1997

• Various roles in drug development and medical affairs.

### ACADEMIC TEACHING & ADMINISTRATIVE EXPERIENCE

**Temple University**Quality Assurance/Regulatory Affairs Program
Adjunct Professor

Philadelphia, PA, USA

2006-Current

### National University of Singapore

Singapore Academy of GxP Excellence (SAGE)

Steering Committee Member

### Singapore

2008

### Monash Medical Center

Melbourne, Victoria, Australia

### Senior Medical Staff: Hypertension & Vascular Medicine

1995-1997

- Investigation, diagnosis and management of complex hypertension patients in the outpatient and inpatient setting.
- Research Fellow, antioxidant research and epidemiology research, Public Health Research and Development Fellow (PH&RDC of NH&MRC).
- Assessor, Quality Indicators in Hospital Care (QUALITY) Study.
- Research Fellow, Department Epidemiology & Preventive Medicine.

# Monash University Alfred Medical School

Melbourne, Victoria, Australia

1993-1997

- Associate Sub-Dean Faculty of Medicine
- Lecturer and coordinator, fifth-year medicine teaching program
- Lecturer and tutor in Epidemiology, Prevention Medicine and Bio-Statistics

### National Health & Medical Research Council Research Scholar

Melbourne, Victoria, Australia

1993-1997

Department of Epidemiology & Preventive Medicine

Melbourne, Victoria, Australia

### Sub-Dean, Austin Hospital Clinical School

Melbourne University Private

1992-1994

- Designed, implemented and administered clinical training program for 300+ third- through sixth-year medical students based at a large specialist tertiary referral hospital.
- Oversaw clinical examinations for medical program and specialist training programs in internal medicine conducted at the Austin hospital.

### Royal Australasian College of Physicians

Melbourne, Victoria, Australia

**Internal Medicine Training Program** 

1991-1994

Vascular medicine & clinical pharmacology

### University of Melbourne

Melbourne, Victoria, Australia

Medical faculty, Clinical Supervisor, Internal Medicine

1989

### **EDUCATION & PROFESSIONAL CREDENTIALS**

University of Melbourne

Melbourne, Victoria, Australia

**Doctor of Medicine (MD) degree** 

1989

Bachelor of Medicine (MBBS) degree

Graduate and Undergraduate Medical Training

University of Melbourne, Medical Faculty

Royal Melbourne Hospital Clinical School

Fellow, Royal Australasian College of Physicians

Australia

Member, Royal Australasian College of Medical Administrators

Australia

### **MILITARY SERVICE**

### Royal Australian Armoured Corps

1980-1983

4<sup>th</sup> /19<sup>th</sup> Prince of Wales Light Horse Regiment B Squadron (Trooper/Gunner/Driver/Signaller M113 A1 Armoured Personnel Carrier) Royal Australian Army Reserve, Officer Training School

### **MISCELLANEOUS**

Lethbridge Wines, a commercial vineyard

1996-Current

Founder/Partner

### **Aspen Institute First Mover Fellow**

2013

### **SELECT PUBLICATIONS & PRESENTATIONS** (abbreviated list)

- Witness, U.S. House of Representatives hearing on "Meeting the Challenge of Drug-Resistant Diseases in Developing Countries." Written testimony delivered on behalf of Johnson & Johnson before the House Subcommittee on Africa, Global Health, Global Human Rights, and International Organizations: Washington, DC, April 2013.
- Panelist, U.S.-China Food and Drug Law Conference, "Innovation and Access: Key Success Factors in China," hosted by the Food & Drug Law Institute, Beijing, June 2011.
- Presenter and panelist, University of Miami Global Business Forum, "Incentives for Innovation: Who's the customer?" January 2011.
- Co-author, "Comparative Effectiveness, Personalized Medicine and Innovation: The Path Forward." PharmacoEconomics, October 2010.
- Contributor, Canadian Roundtable on Clinical Considerations for Subsequent Entry Biologics, "The Global Experience with Subsequent Entry Biologics," May 2009.
- Co-editor, Global Pharmacovigilance Laws and Regulations: The Essential Reference, Food and Drug Law Institute, April 2009.
- Spokesperson for Johnson & Johnson on the topic of the Healthcare Notification Network (HCNN) partnership, October 2008.
- Presenter, Brandeis University Health Industry Forum, "Enhancing Post-Marketing Surveillance," April 2007.
- Co-author, "Pharmacovigilance," Chapter 9 of *The Learning Healthcare System: Workshop Summary*, a publication of the Institute of Medicine, Washington, DC, 2007.
- Invited witness, U.S. Senate hearing on "Building a 21st Century FDA: Proposals to Improve Drug Safety and Innovation." Testimony delivered on behalf of Johnson & Johnson before the Senate Committee on Health, Education, Labor, and Pensions: Washington, DC, November 2006.
- Presenter, "Epidemiology and surveillance in a billion dollar whodunit," Seminar Series, Monash University Department of Epidemiology & Preventive Medicine, Melbourne, Australia, August 2004.

# **Committee on Energy and Commerce**

U.S. House of Representatives
Witness Disclosure Requirement - "Truth in Testimony"
Required by House Rule XI, Clause 2(g)

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1.	Your Name: Adrian Thomas, MD, FRACP	9	
2.	Are you testifying on behalf of the Federal, or a State or local government entity?	Yes	No X
3.	Are you testifying on behalf of an entity that is not a government entity?	Yes X	No
4.	Other than yourself, please list which entity or entities you are representing	ng:	
	Janssen Research and Development, L.L.C		
5.	Please list any Federal grants or contracts (including subgrants or subcontracts you or the entity you represent have received on or after October 1, 2011: See attached		that
6.	If your answer to the question in item 3 in this form is "yes," please descriposition or representational capacity with the entity or entities you are represented and Vice President, Global Market Access & Commercial Strategy Oper Head, Global Public Health	presenti	ng:
7.	If your answer to the question in item 3 is "yes," do any of the entities disclosed in item 4 have parent organizations, subsidiaries, or partnerships that you are not representing in your testimony?	Yes X	No
8.	If the answer to the question in item 3 is "yes," please list any Federal grace contracts (including subgrants or subcontracts) that were received by the under the question in item 4 on or after October 1, 2011, that exceed 10 per revenue of the entities in the year received, including the source and amongrant or contract to be listed:  None	entities ercent o	f the
9.	Please attach your curriculum vitae to your completed disclosure form.	a e	
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Sig	nature:Date:	17-	-2011

Contract Number	Disease Area	Period of Performance	Crucell as subcontractor	Crucell as subcontractor   Primary Contract / Grant holder   Amount Awarded (USD)	Amount Awarded (USD)	Awarding Party
IPCAVD - 1- A1066305	HIV Vaccine Research*	2005-2011	Yes	Harvard University	8.4MM**	NIAID
IPCAVD -2- multivalent A1078526	HIV Vaccine Research*	2008-2013	Yes	Harvard University	27.7MM	NIAID
IPCAVD -3- A1096040	HIV Vaccine Research*	2012-2017	Yes	Harvard University	7.5MM	NIAID
HHSN272200800056C	Multivalent Filovirus Vaccine Development	2008-2015	No	Crucell	13.4MM	NIAID
HHSN272200800056C	Multivalent Filovirus Vaccine Development	2010-2015	No	Crucell	17.4MM	NIAID
HHSN272200800056C	Multivalent Filovirus Vaccine Development	2010-2015	No N	Crucell	3.5MM	NIAID
N01-A1-05421	Malaria Vaccine Research	2008-2011	Yes	SAIC	2.9MM	NIAID
Under LOI (MVI-Crucell)	Malaria Vaccine Research	2010-2012	Yes	MVI	2.0MM	USAID
HHSN272200900060C	Infectious Disease - Respiratory - Flu A	2009-2014	No	Crucell	47.0MM	NIAID

\*not a "neglected" disease, but included due to its profound importance in many developing countries \*\*Much, probably over half, spent prior to the period in question