

### www.isctr.org Translational Pathways for Cardiovascular Devices - Online Course -

# 80 Multidisciplinary Lectures presented by Innovators, Industry, Regulatory (FDA & EU), Reimbursement, Practice Guideline, and Patients Target Audience:

Inventors, Clinical and Basic Scientists, Interventional Cardiologists, Medical Students, Engineers, Industry, Regulators, Payers, and Investors

Introduction (Total 12.5 mins)

1) Welcome from the Course Directors (View for free online)

- Spencer King, MD, Emory University (5 mins 50 secs)
- Anthony DeMaria, MD, University of California San Diego (5 mins)
- Nabil Dib, MD, ISCTR (1 min 40 secs)

Session I: Basic Knowledge for CV Devices Development (Total = 10 hrs)

## **Topic 1: Concept/Innovation** (*Total = 7 mins*)

- 1) Choosing an Innovative Concept
  - Todd Brinton, MD, Edwards Lifesciences (7 mins 10 secs)

### **Topic 2: Intellectual Property** (*Total = 27 mins*)

#### 2) Intellectual Property

- James Inskeep, Inskeep Intellectual Property Group (4 mins 45 secs)
- 3) Overview of the Patent Process
  - Jason Gilbert, Inskeep Intellectual Property Group (22 mins)

## **Topic 3: Business Plan, Product Development, and Fundraising** (*Total = 6 mins*)

## 4) Business Plan, Product Development, and Fundraising

• Stan Rowe, NXT Biomedical (6 mins)

## **Topic 4: Product Manufacturing** (*Total = 24 mins*)

- 5) **Requirements for Medical Device Manufacturing & Iteration FDA Point of View** 
  - Brad Quinn, FDA (9 mins 35 secs)
- 6) Requirements for Medical Device Manufacturing & Iteration Industry Point of View
  - Richard Rapoza, PhD, Abbott Vascular (14 mins 35 secs)

**Topic 5: Preclinical Evaluation/Animal Model** (*Total = 1 hr*)

- 7) Advanced Cardiac Anatomy Application in Translational Research Tailored to Current and Future Technology
  - Renu Virmani, MD, CV Path Institute (14 mins 10 secs)
- 8) Large Animal Model for Heart Failure, Valvular Disease, Coronary Artery Disease, and Device Testing
  - Daniel Burkhoff, MD, Columbia University (15 mins 50 secs)
- 9) Pre-Clinical Study Design & Endpoints for Device Evaluation FDA Point of View
  Judith Davis, DVM, MS, FDA (16 mins)

### 10) Pre-Clinical Study Design & Endpoints for Device Evaluation – Investigator Point of View

• Renu Virmani, MD, CV Path Institute (12 mins 40 secs)

**Topic 6: Early Feasibility** (*Total = 31 mins*)

#### 11) Early Feasibility Studies for Device Evaluation

- Andrew Farb, MD, FDA (14 mins 50 secs)
- 12) Current Challenges & Future Direction for Human Early Feasibility Study for Device Evaluation – Industry Point of View
  - David Reuter, MD, Seattle Children's Hospital (15 mins 35 secs)

### **Topic 7: Biostatistics** (Total = 2 hrs 22 mins)

- 13) Basic in Statistics Clinical Study Design for Translational Research
  - Chris Mullin, PhD, NAMSA (16 mins)
- 14) Basic Statistical Concepts
  - Chris Mullin, PhD, NAMSA (22 mins 25 secs)
- 15) Sample Size and Power
  - Chris Mullin, PhD, NAMSA (22 mins 30 secs)
- 16) Sensitivity and Specificity
  - Chris Mullin, PhD, NAMSA (9 mins 15 secs)
- 17) Common Study Design
  - Chris Mullin, PhD, NAMSA (22 mins 50 secs)
- 18) Phases of Translational Research
  - Chris Mullin, PhD, NAMSA (4 mins 15 secs)
- 19) Statistics for Evaluation of Cardiovascular Diagnostic Devices
  - Chris Mullin, PhD, NAMSA (12 mins 40 secs)
- 20) **Pre-Clinical & Clinical Trial Design & Endpoints of Fast Track to Device Approval** 
  - Roseann White, PhD, Duke Research Institute (14 mins 45 secs)
- 21) Advanced Statistical Methods for Translational Research
  - Chris Mullin, PhD, NAMSA (13 mins 45 secs)
- 22) Clinical Endpoints/Surrogate Endpoints
  - Roseann White, PhD, Duke Research Institute (6 mins 15 secs)

## Topic 8: Regulatory Approval (Total = 48 mins)

### 23) Regulatory Requirement for Marketing Approval

• Bram Zuckerman, MD, FDA (View for free online) (15 mins 20 secs)

### 24) FDA Perspective on Transformative Regulatory Pathways & Device Innovation

- Bram Zuckerman, MD, FDA (9 mins 10 secs)
- 25) Regulatory Review of Cardiovascular Diagnostic Devices FDA Perspective
  - Marco Cannella, PhD, FDA (9 mins 45 secs)
- 26) Regulatory Review of Cardiovascular Devices European Regulatory Perspective
  - Robert Byrne, MD, Heart Center, Germany (14 mins 10 secs)

## **Topic 9: Reimbursement** (Total = 24 mins)

#### 27) CMS Criteria for Reimbursement for Cardiovascular Innovation

• Joseph Chin, MD, Centers for Medicare and Medicaid Services (CMS) (13 mins 25 secs)

#### 28) Reimbursement for Diagnostic Devices

• Lori Ashby, Centers for Medicare and Medicaid Services (CMS) (10 mins 35 secs)

### Topic 10: Practice Guideline (Total = 32 mins)

#### 29) Practice Guideline Requirement for New Technology

• Alice Jacobs, MD, Boston University (16 mins 20 secs)

#### 30) Guideline Requirements for Diagnostic Devices

• Roxana Mehran, MD, Icahn School of Medicine, Mount Sinai (16 mins)

### **Topic 11: Technology Adoption** (Total = 37 mins)

#### 31) Adoption of Technology

• Ian Meredith, MD, Boston Scientific (7 mins 10 secs)

#### 32) Global Heart Health, Implications for Translational Research

• Salim Yusuf, MD, World Federation of Cardiology (29 mins 25 secs)

## **Topic 12: Conflict of Interest** (Total = 8 mins)

#### 33) Conflict of Interest and Product Development

• Anthony DeMaria, MD, University of California San Diego (7 mins 45 secs)

## **Topic 13: Patients** (Total = 8 mins)

### 34) The Patients Voice

• Mark Mercola, PhD, Stanford Cardiovascular Institute (7 mins 35 secs)

#### **Panel Discussions** (Total = 2 hrs 3 mins)

- Diagnostic Devices Development (32 mins)
- Device Development: Essential Concepts (21 mins)
- Requirements for CV Devices Approval (36 mins 25 secs)
- Clinical Endpoints & Biostatistics (19 mins 30 secs)
- Preclinical, Early Feasibility, and Safety Study for Device Development (14 mins 20 secs)

# Session II: Translational Pathway for Transcatheter Aortic Valve Replacement (Total = 2 hrs 30 mins)

- 1) The Clinical Need for Innovative Treatment for Aortic Valve Disease
  - Martin Leon, MD, Columbia University (17 mins 10 secs)
- 2) The Methods for TAVR Development
  - Stan Rowe, Edwards Lifesciences (10 mins 30 secs)
- 3) The Endpoints for TAVR Development
  - Ori Ben-Yehuda, MD, Cardiovascular Research Foundation (14 mins)
- 4) Current Challenges & Future Direction for AV Development & Iteration FDA Point of View
  Nicole Ibrahim, PhD, FDA (11 mins 30 secs)
- 5) Current Challenges & Future Direction for AV Development & Iteration Industry Point of View
  - Stan Rowe, Edwards Lifesciences (9 mins 15 secs)
- 6) TAVR Development from Concept to First In Man
  - Alain Cribier, MD, University of Rouen, France (View for free online) (18 mins)
- 7) TAVR Development from First In Man to Phase 3 and Beyond
  - Martin Leon, MD, Columbia University (View for free online) (24 mins)

**Panel Discussions** (Total = 46 mins)

- Valve Disease/TAVR (29 mins)
- Aortic Valve Development (16 mins 40 secs)

Session III: Translational Pathway for Transcatheter Mitral/Tricuspid Valve Devices (Total = 2 hrs 38 mins)

- 1) The Clinical Need for Innovative Treatment for Mitral/Tricuspid Valve Disease
  - Michael Mack, MD, Baylor Scott & White Health (15 mins)
- 2) The Methods for Translational Mitra/Tricuspid Valve Device Development
  - Michael Mack, MD, Baylor Scott & White Health (15 mins)
- 3) The Endpoints for Transcatheter Mitral/Tricuspid Valve Device Development
  Blasé Carabello, MD, East Carolina University (12 mins 40 secs)
- 4) Current Challenges & Future Direction for Mitral/Tricuspid Valve Device Development & Iteration FDA Point of View
  - John Laschinger, MD, Edwards Lifesciences (14 mins 45 secs)
- 5) Current Challenges & Future Direction for Mitral/Tricuspid Valve Device Development & Iteration Industry Point of View
  - Patricia Todd, Edwards Lifesciences (14 mins 15 secs)
- 6) Unmet Clinical Needs for Tricuspid Valve Interventions
  - Carlos Sanchez, MD, Ohio Health-Riverside Methodist Hospital (7 mins)
- 7) Current Imaging Limitations for the Advancement of Tricuspid Valve Interventions
   Rebecca Hahn, MD, Columbia University Medical Center (8 mins)
- 8) Tricuspid Valve Interventions: Challenges from the Regulatory Perspective
  - Changfu Wu, PhD, FDA (7 mins)

#### **Panel Discussions** (Total = 1 hr 4 mins)

- Structural Heart Intervention Imaging (15 mins 15 secs)
- MV/TV Transcatheter Repair/Replacement (35 mins 25 secs)
- MV/TV Development (13 mins 20 secs)

# **Session IV: Translational Pathway for Coronary Stent** (*Total* = 3.5 hrs)

- 1) The Clinical Need for Innovative Coronary Stent
  - Gregg Stone, MD, Icahn School of Medicine, Mount Sinai (16 mins 40 secs)
- 2) The Methods for Coronary Stent Development
  - Chuck Simonton, MD, Abiomed (24 mins)
- 3) The Endpoints for Coronary Stent Development
  - Donald Cutlip, MD, Beth Israel-Deaconess Medical Center (10 mins 20 secs)
- 4) Current Challenges & Future Direction for Coronary Stent Development & Iteration FDA Point of View
  - Michael John, MPH, FDA (10 mins)
- 5) Current Challenges & Future Direction for Coronary Stent Development & Iteration Industry Point of View
  - Chuck Simonton, MD, Abiomed (15 mins 30 secs)
- 6) Unmet Clinical Needs, Value Added & Future Direction in CT Lesion Assessment
  - James Min, MD, Weill Cornell Medicine (7 mins)
- 7) Unmet Clinical Needs, Current & Future Direction in Intracoronary Physiology & Imaging Assessment
  - Morton Kern, MD, University of California Irvine & Long Beach Veterans Administration Medical Center (8 mins 20 secs)
- 8) Advances in the Assessment of High-Risk Coronary Lesions Non-Clinical Evaluation
  - Robert Safian, MD, Center for Innovation & Research in CV Diseases (CIRC)(7 mins 30 secs)
- 9) Advances in the Assessment of High-Risk Coronary Lesions FDA Perspective
  - Shawn Forrest, FDA (6 mins 45 secs)
- 10) Revascularization & Devices for Complex Coronary Lesions Calcified & Total Occlusions - Unmet Clinical Needs/Future Directions
  - Ajay Kirtane, MD, Columbia University Medical Center (6 mins 50 secs)
- 11) Revascularization & Devices for Complex Coronary Lesions Calcified & Total Occlusions Non-Clinical Evaluation
  - Kevin Croce, MD, PhD, Harvard Medical School (7 min 40 secs)
- 12) Revascularization & Devices for Complex Coronary Lesions Calcified & Total Occlusions - FDA Perspective
  - Lydia Glaw, PhD, FDA (6 min 15 secs)

**Panel Discussions** (Total = 1 hr 15 mins)

- Devices for Complex Coronary Lesions (17 mins 15 secs)
- Advances in the Assessment of High Risk Coronary Lesions (17 mins 40 secs)

- CAD/Coronary Stent (27 mins)
- Coronary Stent Development (15 mins 35 secs)

**Session V: Translational Pathway for Catheter Ablation** (*Total = 2 hrs*)

- 1) The Clinical Need for the Treatment of Arrhythmia Innovative Catheter Ablation
  - Douglas Packer, MD, Mayo Clinic (16 min 40 secs)
- 2) Methods for Catheter Ablation Development
  - Douglas Packer, MD, Mayo Clinic (17 mins 50 secs)
- 3) The Endpoints for Catheter Ablation Development
  - Marco Cannella, PhD, FDA (6 mins 45 secs)
- 4) Current Challenges & Future Direction for Catheter Ablation Development & Iteration - FDA Point of View
  - Mark Fellman, MS, FDA (14 mins)
- 5) Current Challenges & Future Direction for Catheter Ablation Development & Iteration Industry Point of View
  - Uri Yaron, PhD, Biosense Webster at Johnson & Johnson (14 mins)

#### **Panel Discussions** (*Total = 52 mins*)

- Arrhythmia/Catheter Ablation (29 mins)
- Catheter Ablation Development (23 mins)

Session VI: Translational Pathway for Ventricular Assist Devices (Total = 1 hr 42 mins)

1) Ventricular Assist Devices, the Windy Road to Recovery

- Sir Magdi Yacoub, MD, Imperial College, England (17 mins)
- 2) The Methods for Left Ventricular Assist Devices Development
  - Francis Pagani, MD, PhD, University of Michigan (19 mins)

#### 3) The Endpoints for Left Ventricular Device Evaluation

• Keith Aaronson, MD, University of Michigan (8 mins 25 secs)

**Panel Discussions** (Total = 58 mins)

- Translational Pathway for LV Assist Devices (43 mins 45 secs)
- CHF/Ventricular Assist Devices (14 min 10 secs)

Session VII: Translational Pathway for Interventional Devices for Heart Failure (Total = 53 mins)

- 1) **Overview of Interventional Devices for Heart Failure** 
  - William Abraham, MD, Ohio State University (7 mins)
- 2) Current Landscape & Future Direction Percutaneous Ventricular Assist Devices
  William O'Neill, MD, Henry Ford Hospital (7 mins 25 secs)
- 3) Current Landscape & Future Direction of Neuromodulation Heart Failure Therapies
  - Horst Sievert, MD, CardioVascular Center, Germany (6 mins 40 secs)

- 4) Current Landscape & Future Direction Intracardiac Shunts & Ventricular Remodeling Therapies
  - Gregg Stone, MD, Icahn School of Medicine, Mount Sinai (7 mins 15 secs)
- 5) Interventional Devices for Heart Failure Non-Clinical Evaluation
  - Navin Kapur, MD, Tufts Medical Center (7 mins)
- 6) Interventional Devices for Heart Failure FDA Perspective
  - Ileana Piña, MD, Central Michigan University (7 mins)

**Panel Discussion** (*Total = 11 mins*)

• Interventional Devices for Heart Failure (11 mins)

Session VIII: Translational Pathway for Left Atrial Appendage Closure Devices (Total = 43 mins)

- 1) Unmet Clinical Needs/Current and Future Direction
  - Brian Whisenant, MD, University of Utah (7 mins)
- 2) Current & Future Left Atrial Appendage Imaging Modalities to Optimize LAA Closure
  - Dee Dee Wang, MD, Henry Ford Hospital (6 mins 45 secs)
- 3) Left Atrial Appendage Closure Devices FDA Perspective
  - Rachel Neubrander, PhD, FDA (6 mins 25 secs)

**Panel Discussion** (Total = 23 mins)

• Left Atrial Appendage Closure Devices (22 mins 35 secs)

# **Total Course Hours = approx. 25 hours of lectures**

**Supplemental Lectures in Cardiovascular Medicine** (Total = 1 hr 38 mins)

### 1) Introduction to the Cardiac Cath Lab

- Morton Kern, MD, University of California Irvine & Long Beach Veterans Administration Medical Center (52 mins)
- 2) Cardiac Imaging: The Right Test for the Right Patient
  - Anthony DeMaria, MD, University of California San Diego (46 mins)