

6.12.2019

THE EU MEDICAL DEVICES REGULATION. IS THE MEDTECH ECOSYSTEM READY?

AULA 01 - H. 10.30 - 16.00
VIA BOCCONI, 8 - MILANO

PRIVILEGED TALKS

EXCLUSIVE CONVERSATIONS WITH TOP EUROPEAN POLICY MAKERS

10.30 WELCOME COFFEE & REGISTRATION

11.00 INTRODUCTION TO THE PRIVILEGED TALK
Rosanna Tarricone, Associate Dean Government,
Health and Nonprofit, SDA Bocconi

PART 1 THE EU MEDICAL DEVICE REGULATION
CHAIR: *Aleksandra Torbica*, CERGAS Director,
SDA Bocconi

11.30 THE EU MEDICAL DEVICE REGULATIONS:
INTRODUCTION AND IMPLEMENTATION
Salvatore D'Acunto, Head of the Health
Technology and Cosmetics Unit at European
Commission
Salvatore Scalzo, Policy and Legal Officer at
European Commission

12.30 EVIDENCE GENERATION THROUGHOUT THE
LIFECYCLE OF MDs
Oriana Ciani, Associate Professor, SDA Bocconi

12.50 Q&A

13.15 LUNCH BREAK

PART 2 CHALLENGES AND IMPACTS

CHAIR: *Giuditta Callea*, Associate Professor,
SDA Bocconi

14.00 HOW IS FDA SUPPORTING INNOVATION AND
FACILITATING REGULATORY ALIGNMENT?
Ritu Nalubola, Director, FDA Europe Office

14.15 IS THE MEDTECH INDUSTRY UP TO NEW
CHALLENGES?
Dario Pirovano, Senior Regulatory Adviser
MedTech Europe

14.30 MDR & SAFETY FOR THE PATIENTS. IS THAT
ALL WE NEED?
Valentina Strammiello, Senior Programme
Manager of European Patients' Forum

14.45 Q&A

15.30 CONCLUSION AND RECOMMENDATIONS
Rosanna Tarricone, Associate Dean Government,
Health and Nonprofit, SDA Bocconi