

Con il patrocinio di

**SIE** - Società Italiana di Ematologia



**Lucca, 15<sup>th</sup>-16<sup>th</sup> June 2016**

**IMT Institute for advanced studies**

**Piazza S. Francesco, 19**

**PROGRAMMA  
PRELIMINARE**

**Wednesday, 15<sup>th</sup> June 2016**

### The Young Biomarkers Day

10.00 - 10.30	Welcome Director of IMT School for Advanced Studies Lucca Rector, University of Pisa General Director, University Hospital Director of Department of Clinical and Experimental Medicine Director of Department of Laboratory Medicine	<i>P. Pietrini M. M. Augello C. R. Tomassini M. Petrini R. Danesi</i>
	<b>Chairmen: S. Galimberti &amp; R. Danesi</b>	
10.30 - 11.00	PGx data: clinical application in oncology	<i>R. van Schaik</i>
11.00 - 11.30	Pharmacokinetic modeling incorporating PGx data: clinical application in hematology	<i>A. Di Paolo</i>
11.30 - 12.00	Coffee break	
	<b>Chairmen: M. Del Re &amp; F. Massari</b>	
12.00 - 12.30	Clinical trials for the study of biomarkers: from classical to innovative designs	<i>M. Di Maio</i>
12.30 - 13.00	Clinical trial design: how to validate circulating nucleic acids?	<i>S. Pilotto</i>
13.00 - 14.30	Lunch	
	<b>Chairmen: M. Di Maio &amp; C. Cremolini</b>	
14.30 - 15.00	Clinical case I – CNAs in CRC	<i>C. Antoniotti</i>
15.00 - 15.30	Clinical case II – CNAs in NSCLC	<i>I. Petrini</i>
15.30 - 16.00	Clinical case III – CNAs in mBC	<i>M. Lambertini</i>
16.00 - 16.30	Coffee break	
	<b>Incorporating PGx into clinical practice</b>	
16.30 - 17.00	The role of the pharmacologist	<i>E. Dreussi</i>
17.00 - 17.30	The interest of the oncologist	<i>E. Vasile</i>

**Thursday, 16<sup>th</sup> June 2016**

### Golden Helix Pharmacogenomics Day

09.00 - 09.30	Welcome	<i>R. Danesi, G. Patrinos</i>
	<b>Chairmen: F. Citterio &amp; U. Boggi</b>	
09.30 - 10.00	Pharmacogenomics in the era of Precision Medicine	<i>G. Patrinos</i>
10.00 - 10.30	How and when to monitor CNAs in transplantation	<i>R. van Schaik</i>
10.30 - 11.00	Clinical applications of CNAs in renal transplant	<i>D. A. Hesselink</i>
11.00 - 11.30	Coffee break	
	<b>Chairmen: G. Fontanini &amp; A. Falcone</b>	
11.30 - 12.00	How and when to monitor CNAs in oncology	<i>R. Danesi</i>
12.30 - 12.30	Pros and cons of tissue vs CNAs	<i>A. Scarpa</i>
12.30 - 12.50	The future of Personalize Health Care: The Biotech experience	<i>A. Kiermaier/R. Scalomagna</i>
12.50 - 14.30	Lunch	
	<b>Chairmen: R. Rosell &amp; G. Masi</b>	
14.30 - 15.00	Clinical applications of CNAs in NSCLC	<i>N. Karachaliou</i>
15.00 - 15.30	Clinical applications of CNAs in colorectal cancer	<i>C. Cremolini</i>
15.30 - 16.00	Clinical applications of CNAs in breast cancer	<i>I. Garcia-Murillas</i>
16.00 - 16.30	Coffee break	
	<b>Chairmen: R. van Schaik &amp; S. Galimberti</b>	
16.30 - 17.00	Clinical applications of CNAs in prostate cancer	<i>G. Jenster</i>
17.00 - 17.30	Clinical applications of CNAs in haematologic malignancies	<i>S. Ferrero</i>
17.30 - 18.00	Are we ready for prime time use?	<i>G. Argilés Martínez</i>
18.00 - 18.30	Conclusions and quiz ECM	

## Scientific Directors

Romano Danesi  
George Patrinos  
Marzia Del Re

## Scientific Secretariat

Chiara Cremolini  
Francesco Massari

## Faculty

Guillem Argilés Martínez  
Carlotta Antoniotti  
Massimo Mario Augello  
Ugo Boggi  
Franco Citterio  
Chiara Cremolini  
Romano Danesi  
Marzia Del Re  
Massimo Di Maio

Antonello Di Paolo  
Eva Dreussi  
Alfredo Falcone  
Simone Ferrero  
Gabriella Fontanini  
Sara Galimberti  
Isaac Garcia-Murillas  
Dennis Hesselink  
Guido Jenster

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Iacopo Petrini  
Mario Petrini  
Pietro Pietrini

Sara Pilotto  
Rafael Rosell  
Roberto Scalamogna  
Aldo Scarpa  
Carlo Rinaldo Tomassini  
Ron van Schaik  
Enrico Vasile

## Rationale YBD-GHF

The search for biomarkers is increasingly required in many disciplines: in cancer, to assess the response to treatment and risk of treatment-related toxicity, as well as in transplants to monitor the graft injury. Significant achievements have been made in our understanding and clinical application of pharmacogenetics in cancer of drug-metabolising enzymes in order to predict fluoropyrimidine- and irinotecan-related toxicities with the screening of DPYD and UGT genes, respectively. Circulating tumour DNA could represent a powerful diagnostic tool to help clinical decision making, as it is being used, at least for research purposes, to assess tumour heterogeneity, identify genetic determinants for targeted therapy, evaluate the response to treatment and assess the evolution of molecular resistance over time. Moreover, the field of solid organ transplantation has been seeing improved graft outcomes, with reduced rates of acute rejection due to the use of immunosuppressive drugs (tacrolimus, cyclosporine, MMF). However, the diagnostic tools available to monitor the transplant rejection are costly, have the danger of complications or are available too late for the patient. A new and innovative approach for detecting organ rejection makes use of plasma graft-derived cell free DNA in the recipient, arising from damaged cells when the transplanted donor organ is subject to rejection.

However, the choice of genetic markers and the implementation of new technologies in clinical practice should be discussed between the main stakeholders.

The Young Biomarkers Day and the Golden Helix Symposium on Circulating Nucleic Acids will address these important issues by involving young researchers and expert scientists, with the aim of discussing the appropriate use of technological platforms and molecular biomarkers in transplants and in solid tumours (i.e. colorectal, lung, breast cancer and melanoma), especially in the era of new therapeutic options (i.e. immunotherapy, monoclonal antibodies and new immunosuppressive drugs).

## Informazioni ECM

- Data 15/05/2016 evento nr: 158540
- Data 16/05/2016 evento nr: 158544
- Evento accreditato per:
  - Professione: Medico Chirurgo (tutte le discipline), Biologo, Farmacista, Infermiere, Tecnico Sanitario Laboratorio Biomedico
- Obiettivo formativo: Contenuti tecnico-professionali (conoscenze e competenze) specifici di ciascuna professione, di ciascuna specializzazione e di ciascuna attività ultraspecialistica. Malattie rare
- Ai fini dell'acquisizione dei crediti formativi è necessaria la presenza effettiva per il 100% della durata complessiva dell'evento ed il superamento delle verifiche di apprendimento
- Il limite massimo dei crediti formativi ricondotti al triennio di riferimento acquisibili mediante reclutamento diretto è di 1/3

## Provider ECM - 316

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