

PLATINUM SPONSOR

SANOFI GENZYME 



Shire è ora parte di Takeda

GOLD SPONSOR



SILVER SPONSOR



BRONZE SPONSOR



OTHERS

Baxter

Medtronic
Further Together

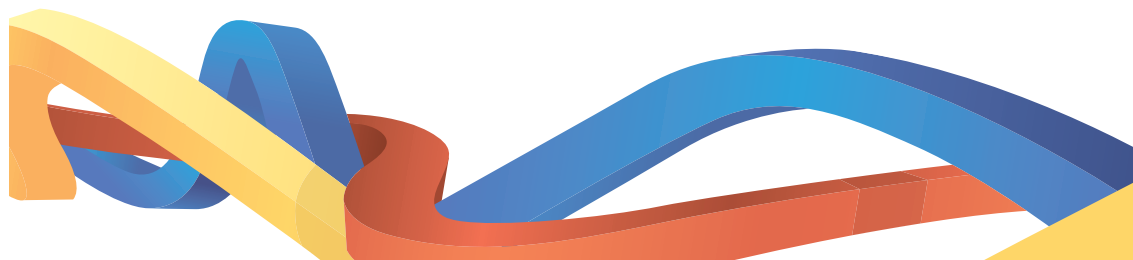


UNIVERSITÀ
DEGLI STUDI
DI BRESCIA

INTERNATIONAL MEETING
**GENETICS OF KIDNEY
DISEASE IN 2019**

Joint Meeting with Columbia University, New York

BRESCIA, ITALY, NOVEMBER 22/23 2019



<u>F. Alberici</u>	Milano	<u>R. Magistroni</u>	Modena
<u>A. Amoroso</u>	Torino	<u>C. Marcantoni</u>	Catania
<u>E. Ars</u>	Barcelona, Spain	<u>R.U. Muller</u>	Cologne, Germany
<u>A. Boletta</u>	Milano	<u>A. Ong</u>	Sheffield, UK
<u>G. Boscutti</u>	Udine	<u>A. Pani</u>	Cagliari
<u>G. Capasso</u>	Napoli	<u>Y. Pei</u>	Toronto, Canada
<u>L. Carlassara</u>	Vicenza	<u>G. Piscopo</u>	Bari
<u>G. Carraro</u>	Padova	<u>C. Ponticelli</u>	Milano
<u>L. Cirami</u>	Firenze	<u>F. Prefumo</u>	Brescia
<u>E. Daina</u>	Bergamo	<u>L. Rampoldi</u>	Milano
<u>N. Dalleria</u>	Brescia	<u>S. Sanna Cherchi</u>	New York, USA
<u>E. Delbarba</u>	Brescia	<u>D. Santoro</u>	Messina
<u>L. Econimo</u>	Brescia	<u>G. Savoldi</u>	Brescia
<u>S. Feriozzi</u>	Viterbo	<u>S. Savoldi</u>	Torino
<u>M. Gennarelli</u>	Brescia	<u>F. Scolari</u>	Brescia
<u>P.D. Germain</u>	Paris, France	<u>L. Sternfeld Pavia</u>	Milano
<u>L. Gesualdo</u>	Bari	<u>R. Torra</u>	Barcelona, Spain
<u>A. Gharavi</u>	New York, USA	<u>F. Torri</u>	Brescia
<u>G. Ghiggeri</u>	Genova	<u>E.M. Valente</u>	Pavia
<u>C. Izzi</u>	Brescia	<u>D. Zuccarello</u>	Padova
<u>K. Kiryluk</u>	New York, USA		

13.30 Welcome coffee

14.00 **OPENING MESSAGE AND WELCOME OF THE AUTHORITIES**

SESSION I - GENETICS OF GLOMERULAR DISEASES

Chairmen: C. Ponticelli, F. Scolari

14.30 **Genetic architecture of membranous nephropathy**
K. Kiryluk

15.00 **Genetics of MCD/FSGS**
S. Sanna Cherchi

15.30 **Genetic association study of kidney allograft rejection**
K. Kiryluk

16.00 **Redefining ANCA-vasculitis: from genetics to treatment through phenotype**
F. Alberici

Chairmen: A. Amoroso, D. Santoro

16.30 **C3 nephropathy: the Italian Registry**
E. Daina

16.50 **Alport syndrome is common but often unrecognized**
R. Torra

17.15 **Genetic testing in suspected monogenic glomerulopathies**
E. Ars

17.35 **Disease activity in long-standing glomerular disease**
E. Delbarba

17.45 **Family history of nephropathy and disease characteristics in primary glomerular disease**
L. Carlassara

18.10 **MAGISTRAL LECTURE**
Chairmen: C. Izzi, F. Scolari
Towards precision nephrology: opportunities and challenges
A. Gharavi

18.50 End of the Session

SESSION II - HEREDITARY TUBULOINTERSTITIAL KIDNEY DISEASE

Chairmen: M. Gennarelli, F. Prefumo, F. Torri

- 09.00 **Autosomal Dominant Tubulointerstitial Kidney (ADTKD): gene defects and pathophysiology**
L. Rampoldi
- 09.30 **Genetic basis of congenital anomalies of kidney and urinary tract**
S. Sanna Cherchi
- 10.00 **Ciliopathies and kidney**
E.M. Valente
- 10.30 Coffee break

SESSION III - ROUND TABLE ON FABRY'S DISEASE

Chairmen: G. Ghiggeri, A. Pani

- 10.45 **Fabry disease: the great masquerader**
D. Germain
- 11.15 **Whom to treat and when to begin**
G. Carraro
- 11.30 **ERT therapy: alfa-agalsidasi**
S. Feriozzi
- 11.45 **ERT therapy: beta-agalsidasi**
L. Econimo
- 12.00 **Chaperone therapy: Migalastat**
L. Cirami
- 12.15 **Discussion**
- 12.30 Light lunch

SESSION IV - AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE (ADPKD)

Chairmen: L. Gesualdo, C. Marcantoni

- 13.30 **Metabolic alterations in ADPKD**
A. Boletta
- 13.55 **The genetic complexity of ADPKD**
C. Izzi
- 14.20 **Preimplantation Genetic Diagnosis (PGD) in ADPKD**
D. Zuccarello
- 14.45 **Why a PKD clinic?**
A. Ong
- 15.10 **Italian Registry of ADPKD**
C. Izzi
- Chairmen: G. Boscutti, G. Capasso
- 15.30 **Assessment of risk for ADPKD progression**
Y. Pei
- 16.00 **Tolvaptan in real-life clinical care**
R.U. Muller
- 16.30 **Discussion**

NOT C.M.E. SESSION

- 16.45 **10° Workshop Associazione Italiana Rene Policistico onlus**
Chairmen: F. Scolari, L. Sternfeld Pavia
The ADPKD patients meet the experts
A. Boletta, N. Dallerà, C. Izzi, R. Magistroni,
R.U. Muller, A. Ong, Y. Pei, G. Piscopo,
G. Savoldi, S. Savoldi, D. Zuccarello
- 18.30 End of the Meeting



GENERAL INFORMATION

CONGRESS VENUE

Università degli Studi di Brescia
Dipartimento di Medicina
Viale Europa, 11 • 25123 Brescia

CONGRESS PRESIDENT

F. Scolari (Università degli Studi Brescia)

SCIENTIFIC COMMITTEE

A. Gharavi (Columbia University, New York, USA)
C. Izzi (ASST Spedali Civili di Brescia)
K. Kiryluk (Columbia University, New York, USA)
L. Rampoldi (Ospedale San Raffaele, Milano)
S. Sanna Cherchi (Columbia University, New York, USA)

REGISTRATION

For registration visit www.aristea.com/nefrogenetica

C.M.E. - CONTINUING MEDICAL EDUCATION

Aristea Education (Provider n. 500) has included the Event in the Educational Plan 2019. The Session will be suited for Nephrologist, Cardiology, Medical Genetics, Internal Medicine, Pediatrics and Biologist and provides 10 C.M.E. credits. In order to obtain C.M.E. credits, participants must attend the 100% of the Meeting and submit the filled in evaluation questionnaire and C.M.E.: Verification Form.

The certificate of attendance with the number of C.M.E. assigned credits can be downloaded after 60 days from the date of the Congress directly through the website www.aristeaeducation.it

ORGANIZING SECRETARIAT

 aristea

Via Roma, 10 • 16121 Genova • Italy
Tel. +39 010 553591 • Fax +39 010 5535970
E-mail genova@aristea.com • Web www.aristea.com



GENETICS
OF KIDNEY
DISEASE
IN 2019

